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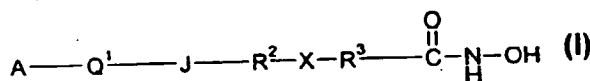
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(54) Title: **CARBAMIC ACID COMPOUNDS COMPRISING AN ETHER LINKAGE AS HDAC INHIBITORS**



amide linkage selected from: $-NR^1C(=O)-$ and $-C(=O)NR^1-$; R_1 is an amido substituent; X is an ether heteroatom, and is $-O-$ or $-S-$; and, R^2 and R^3 are each independently an ether group; and pharmaceutically acceptable salts, solvates, amides, esters, ethers, chemically protected forms, and prodrugs thereof. The present invention also pertains to pharmaceutical compositions comprising such compounds, and the use of such compounds and compositions, both in vitro and in vivo, to inhibit HDAC, and, e.g., to inhibit proliferative conditions, such as cancer and psoriasis.

(57) Abstract: This invention pertains to certain active carbamic acid compounds which inhibit HDAC activity and which have the following formula(I) wherein: A is an aryl group; Q^1 is a covalent bond or an aryl leader group; J is an

CARBAMIC ACID COMPOUNDS COMPRISING AN ETHER LINKAGE AS HDAC INHIBITORS

RELATED APPLICATION

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This application claims priority to United Kingdom Patent Application Number GB 0023983.0 filed 29 September 2000, the contents of which are incorporated herein by reference in their entirety.

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TECHNICAL FIELD

This invention pertains generally to the field of biologically active compounds, and more specifically to certain active carbamic acid compounds which inhibit HDAC (histone deacetylase) activity. The present invention also pertains to pharmaceutical compositions comprising such compounds, and the use of such compounds and compositions, both *in vitro* and *in vivo*, to inhibit HDAC, and, e.g., to inhibit proliferative conditions, such as cancer and psoriasis.

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BACKGROUND

20

DNA in eukaryotic cells is tightly complexed with proteins (histones) to form chromatin. Histones are small, positively charged proteins which are rich in basic amino acids (positively charged at physiological pH), which contact the phosphate groups (negatively charged at physiological pH) of DNA. There are five main classes of histones, H1, H2A, H2B, H3, and H4. The amino acid sequences of histones H2A, H2B, H3, and H4 show remarkable conservation between species, whereas H1 varies somewhat, and in some cases is replaced by another histone, e.g., H5. Four pairs of each of H2A, H2B, H3, and H4 together form a disk-shaped octomeric protein core, around which DNA (about 140 base pairs) is wound to form a nucleosome. Individual nucleosomes are connected by short stretches of linker DNA associated with another histone molecule (e.g., H1, or in certain cases, H5) to form a structure resembling a beaded string, which is itself arranged in a helical stack, known as a solenoid.

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The majority of histones are synthesised during the S phase of the cell cycle, and newly synthesised histones quickly enter the nucleus to become associated with DNA. Within minutes of its synthesis, new DNA becomes associated with
5 histones in nucleosomal structures.

A small fraction of histones, more specifically, the amino side chains thereof, are enzymatically modified by post-translational addition of methyl, acetyl, or phosphate groups, neutralising the positive charge of the side chain, or converting
10 it to a negative charge. For example, lysine and arginine groups may be methylated, lysine groups may be acetylated, and serine groups may be phosphorylated. For lysine, the $-(CH_2)_4-NH_2$ sidechain may be acetylated, for example by an acetyltransferase enzyme, to give the amide
15 $-(CH_2)_4-NHC(=O)CH_3$. Methylation, acetylation, and phosphorylation of amino termini of histones which extend from the nucleosomal core affects chromatin structure and gene expression. (See, for example, Spencer and Davie, 1999).

Acetylation and deacetylation of histones is associated with transcriptional events leading to cell proliferation and/or differentiation. Regulation of the function of
20 transcription factors is also mediated through acetylation. Recent reviews of histone deacetylation include Kouzarides, 1999 and Pazin et al., 1997.

The correlation between the acetylation status of histones and the transcription of genes has been known for over 30 years (see, for example, Howe et al., 1999).
25 Certain enzymes, specifically acetylases (e.g., histone acetyltransferase, HAT) and deacetylases (e.g., histone deacetylase, HDAC), which regulate the acetylation state of histones have been identified in many organisms and have been implicated in the regulation of numerous genes, confirming the link between acetylation and transcription. See, for example, Davie, 1998. In general, histone
30 acetylation correlates with transcriptional activation, whereas histone deacetylation is associated with gene repression.

A growing number of histone deacetylases (HDACs) have been identified (see, for example, Ng and Bird, 2000). The first deacetylase, HDAC1, was identified in 1996 (see, for example, Tauton et al., 1996). Subsequently, two other nuclear mammalian deacetylases has been found, HDAC2 and HDAC3 (see, for example, 5 Yang et al., 1996, 1997, and Emiliani et al., 1998). See also, Grozinger et al., 1999; Kao et al., 2000; and Van den Wyngaert et al., 2000.

Eight human HDACs have been cloned so far:

10 HDAC1 (Genbank Accession No. NP_004955)
HDAC2 (Genbank Accession No. NP_001518)
HDAC3 (Genbank Accession No. O15739)
HDAC4 (Genbank Accession No. AAD29046)
HDAC5 (Genbank Accession No. NP_005465)
HDAC6 (Genbank Accession No. NP_006035)
15 HDAC7 (Genbank Accession No. AAF63491)
HDAC8 (Genbank Accession No. AAF73428)

These eight human HDACs fall in two distinct classes: HDACs 1,2,3 and 8 are in class I, and HDACs 4,5,6 and 7 are in class II.

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There are a number of histone deacetylases in yeast, including the following:

RPD3 (Genbank Accession No. NP_014069)
HDA1 (Genbank Accession No. P53973)
HOS1 (Genbank Accession No. Q12214)
25 HOS2 (Genbank Accession No. P53096)
HOS3 (Genbank Accession No. Q02959)

There are also numerous plant deacetylases, for example, HD2, in *Zea mays* (Genbank Accession No. AF254073_1).

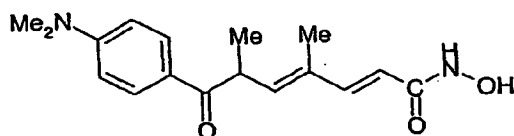
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HDACs function as part of large multiprotein complexes, which are tethered to the promoter and repress transcription. Well characterised transcriptional repressors such as Mad (Laherty et al., 1997), pRb (Brehm et al., 1998), nuclear receptors

(Wong et al., 1998) and YY1 (Yang et al., 1997) associate with HDAC complexes to exert their repressor function.

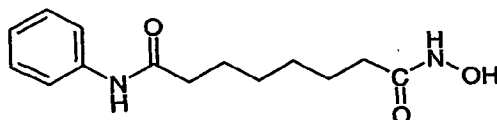
5 The study of inhibitors of histone deacetylases indicates that these enzymes play an important role in cell proliferation and differentiation. The inhibitor Trichostatin A (TSA) (Yoshida et al., 1990a) causes cell cycle arrest at both G1 and G2 phases (Yoshida and Beppu, 1988), reverts the transformed phenotype of different cell lines, and induces differentiation of Friend leukaemia cells and others (Yoshida et al., 1990b). TSA (and SAHA) have been reported to inhibit cell
10 growth, induce terminal differentiation, and prevent the formation of tumours in mice (Finnin et al., 1999).

Trichostatin A (TSA)



15

Suberoylanilide Hydroxamic Acid (SAHA)



20 Cell cycle arrest by TSA correlates with an increased expression of gelsolin (Hoshikawa et al., 1994), an actin regulatory protein that is down regulated in malignant breast cancer (Mielnicki et al., 1999). Similar effects on cell cycle and differentiation have been observed with a number of deacetylase inhibitors (Kim et al., 1999).

25 Trichostatin A has also been reported to be useful in the treatment of fibrosis, e.g., liver fibrosis and liver cirrhosis. See, e.g., Geerts et al., 1998.

Recently, certain compounds that induce differentiation have been reported to inhibit histone deacetylases. Several experimental antitumour compounds, such

- as trichostatin A (TSA), trapoxin, suberoylanilide hydroxamic acid (SAHA), and phenylbutyrate have been reported to act, at least in part, by inhibiting histone deacetylase (see, e.g., Yoshida et al., 1990; Richon et al., 1998; Kijima et al., 1993). Additionally, diallyl sulfide and related molecules (see, e.g., Lea et al., 1999), oxamflatin (see, e.g., Kim et al., 1999), MS-27-275, a synthetic benzamide derivative (see, e.g., Saito et al., 1999; Suzuki et al., 1999; note that MS-27-275 was later re-named as MS-275), butyrate derivatives (see, e.g., Lea and Tulsyan, 1995), FR901228 (see, e.g., Nokajima et al., 1998), depudecin (see, e.g., Kwon et al., 1998), and m-carboxycinnamic acid bishydroxamide (see, e.g., Richon et al., 1998) have been reported to inhibit histone deacetylases. *In vitro*, some of these compounds are reported to inhibit the growth of fibroblast cells by causing cell cycle arrest in the G1 and G2 phases, and can lead to the terminal differentiation and loss of transforming potential of a variety of transformed cell lines (see, e.g., Richon et al., 1996; Kim et al., 1999; Yoshida et al., 1995; Yoshida & Beppu, 1988). *In vivo*, phenylbutyrate is reported to be effective in the treatment of acute promyelocytic leukemia in conjunction with retinoic acid (see, e.g., Warrell et al., 1998). SAHA is reported to be effective in preventing the formation of mammary tumours in rats, and lung tumours in mice (see, e.g., Desai et al., 1999).
- 20 The clear involvement of HDACs in the control of cell proliferation and differentiation suggests that aberrant HDAC activity may play a role in cancer. The most direct demonstration that deacetylases contribute to cancer development comes from the analysis of different acute promyelocytic leukaemias (APL). In most APL patients, a translocation of chromosomes 15 and 17
- 25 (t(15;17)) results in the expression of a fusion protein containing the N-terminal portion of PML gene product linked to most of RAR α (retinoic acid receptor). In some cases, a different translocation (t(11;17)) causes the fusion between the zinc finger protein PLZF and RAR α . In the absence of ligand, the wild type RAR α represses target genes by tethering HDAC repressor complexes to the promoter
- 30 DNA. During normal hematopoiesis, retinoic acid (RA) binds RAR α and displaces the repressor complex, allowing expression of genes implicated in myeloid differentiation. The RAR α fusion proteins occurring in APL patients are no longer responsive to physiological levels of RA and they interfere with the expression of

the RA-inducible genes that promote myeloid differentiation. This results in a clonal expansion of promyelocytic cells and development of leukaemia. *In vitro* experiments have shown that TSA is capable of restoring RA-responsiveness to the fusion RAR α proteins and of allowing myeloid differentiation. These results
5 establish a link between HDACs and oncogenesis and suggest that HDACs are potential targets for pharmaceutical intervention in APL patients. (See, for example, Kitamura et al., 2000; David et al., 1998; Lin et al., 1998).

Furthermore, different lines of evidence suggest that HDACs may be important
10 therapeutic targets in other types of cancer. Cell lines derived from many different cancers (prostate, colorectal, breast, neuronal, hepatic) are induced to differentiate by HDAC inhibitors (Yoshida and Horinouchi, 1999). A number of HDAC inhibitors have been studied in animal models of cancer. They reduce
15 tumour growth and prolong the lifespan of mice bearing different types of transplanted tumours, including melanoma, leukaemia, colon, lung and gastric carcinomas, etc. (Ueda et al., 1994; Kim et al., 1999).

Psoriasis is a common chronic disfiguring skin disease which is characterised by well-demarcated, red, hardened scaly plaques: these may be limited or
20 widespread. The prevalence rate of psoriasis is approximately 2%, i.e., 12.5 million sufferers in the triad countries (US/Europe/Japan). While the disease is rarely fatal, it clearly has serious detrimental effects upon the quality of life of the patient: this is further compounded by the lack of effective therapies. Present
25 treatments are either ineffective, cosmetically unacceptable, or possess undesired side effects. There is therefore a large unmet clinical need for effective and safe drugs for this condition.

Psoriasis is a disease of complex etiology. Whilst there is clearly a genetic component, with a number of gene loci being involved, there are also undefined
30 environmental triggers. Whatever the ultimate cause of psoriasis, at the cellular level, it is characterised by local T-cell mediated inflammation, by keratinocyte hyperproliferation, and by localised angiogenesis. These are all processes in which histone deacetylases have been implicated (see, e.g., Saunders et al.,

1999; Bernhard et al, 1999; Takahashi et al, 1996; Kim et al, 2001). Therefore HDAC inhibitors may be of use in therapy for psoriasis. Candidate drugs may be screened, for example, using proliferation assays with T-cells and/or keratinocytes.

5

Thus, one aim of the present invention is the provision of compounds which are potent inhibitors of histone deacetylases (HDACs). There is a pressing need for such compounds, particularly for use as antiproliferatives, for example, anti-cancer agents, agents for the treatment of psoriasis, etc.

10

Such molecules desirably have one or more of the following properties and/or effects:

- (a) easily gain access to and act upon tumour cells;
- (b) down-regulate HDAC activity;
- 15 (c) inhibit the formation of HDAC complexes;
- (d) inhibit the interactions of HDAC complexes;
- (e) inhibit tumour cell proliferation;
- (e) promote tumour cell apoptosis;
- (f) inhibit tumour growth; and,
- 20 (g) complement the activity of traditional chemotherapeutic agents.

A number of carbamic acid compounds have been described.

Amides

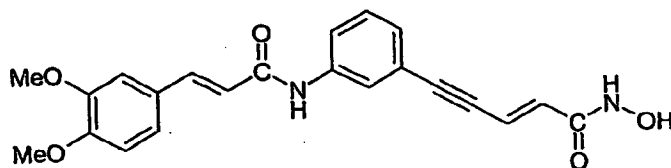
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Hashimoto et al., 1989 describe hydroxamic acid compounds which are claimed to inhibit cell proliferation. Some of the compounds are carbamic acid compounds having a substituted phenyl-dione group linked to a carbamic acid group (-CONHOH) via an aryl-substituted alkylene group.

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Ohtani et al., 1993 describe a number of hydroxamic acid compounds which are claimed to be inhibitors of ras transformation. A few of the compounds are carbamic acid compounds having a phenylacylamido group (-NHCOPh) linked to

a carbamic acid group (-CONHOH) via a phenylene-meta-alkylene group having a carbon-carbon triple bond. See, for example, compounds I-29 (page 69), I-39 (page 87), and I-41 (page 90). Compound I-41, shown below, employs an aryl leader.



5

Onishi et al., 1996, describe several hydroxamic acid compounds which have a phenyl (or substituted phenyl) group linked via an oxazole group to a carbamic acid group. These compounds were reported to inhibit a deacetylase enzyme critical in the biosynthesis of lipid A (a component of the outer membrane of Gram-negative bacteria).

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Parsons et al., 1998 describe a number of hydroxamic acid compounds which are claimed to selectively prevent the growth of a variety of human tumour cell lines.

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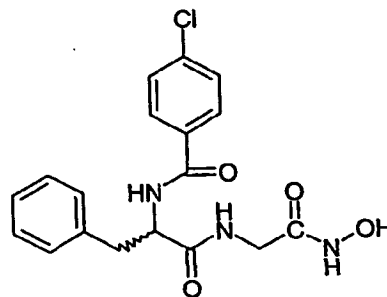
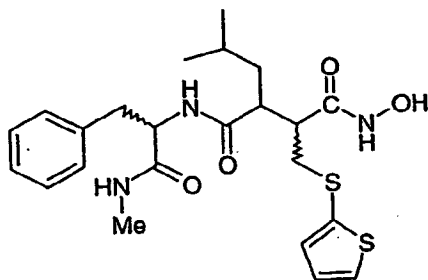
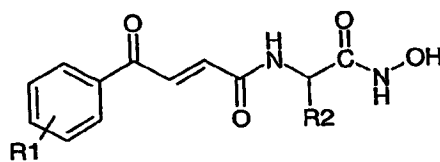
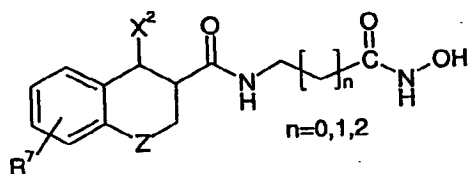
Some of the compounds are carbamic acid compounds having an arylamide group linked to a carbamic acid group via a methylene or substituted methylene group (see, for example, pages 16 and 17).

Some of the compounds are carbamic acid compounds having a phenylamido group (-CONHPh) linked to a carbamic acid group (-CONHOH) via a long alkylene chain, $-(CH_2)_n-$, wherein n is from 4 to 7 (see, for example, pages 47, 48, and 58 therein).

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Some of the compounds are carbamic acid compounds having an aryl group linked via a short chain to an amide group (-CONH-), which in turn is linked via a short chain (e.g., 3 atoms or less) to a carbamic acid group (-CONHOH) (see, for example, page 16, 2nd formula; page 46, 4th formula; page 51, compound 7; and page 61, 2nd formula, therein).

25



Richon et al., 1998 describe several hydroxamic acid compounds, including SAHA, which apparently inhibit HDAC activity, and induce terminal differentiation and/or apoptosis in various transformed cells (see, for example, Table 1 therein).

Suzuki et al., 1998 describe a number of hydroxamic acid compounds which are claimed to have antitumour activity. Some of the compounds are carbamic acid compounds having a substituted phenylamido group (-CONHPh) linked to a carbamic acid (-CONHOH) group via a phenylene-meta-ethenylene or phenylene-para-ethylenylene group (see, for example, pages 8 and 9, compounds 31-50).

Breslow et al., 1994, 1995, 1997 describe a number of hydroxamic acid compounds which are claimed to selectively induce terminal differentiation of neoplastic cells.

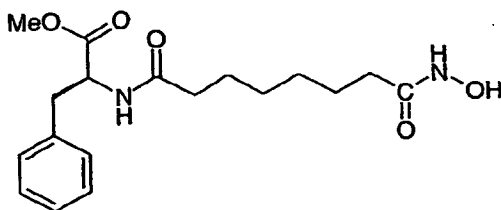
Some of the compounds are carbamic acid compounds having a substituted phenylacylamido group (-NHCOPh) linked to a carbamic acid (-CONHOH) group via a long alkylene chain, $-(CH_2)_n-$, wherein n is from 4 to 8

Some of the compounds are carbamic acid compounds having a substituted phenylamido group (-CONHPh) or phenylacylamido group (-NHCOPh) linked to a carbamic acid (-CONHOH) group via a long alkylene chain, $-(CH_2)_n-$, wherein n is from 4 to 8 (see, for example, columns 7 and 13 of Breslow et al., 1997), or via a

phenylene group (see, for example, columns 24, 30-31 and compounds 20-55 in Table 1 of Breslow et al., 1997).

- One of the compounds is a carbamic acid compound having benzylamido group
5 (-CONHCH₂Ph) linked to a carbamic acid group (-CONHOH) via a -(CH₂)₆- group (see, for example, compound 19 in Table 1, at column 37 of Breslow et al., 1997).

- Jung et al., 1997, 1999, describe several aromatic hydroxamic acid compounds which apparently inhibit HDAC. Some of the compounds have a phenylamido
10 group (PhCONH-). One compound, a peptide analog, is shown below (see, e.g., compound 6 in Jung et al., 1997; compound 4 in Jung et al., 1999).



- Kato et al., 1998, describe a number of aromatic hydroxamic acid compounds,
15 comprising an aryl group linked via an alkylene group to a carbamic acid group, which are apparently active in the treatment of neurodegenerative conditions. One compound, 4-1 at columns 63-64, has a phenylamido group (PhCONH-) linked via a -(CH₂)₅- group to a carbamic acid group.

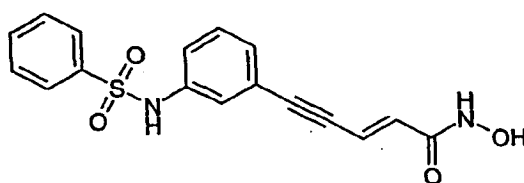
- 20 Glick et al., 1999, describe the apparent apoptotic and differentiating effects of m-carboxy-cinnamic acid bishydroxamide (CBHA) on various tumour cell lines.

- Massa et al., 2001, describe various hydroxamic acid compounds which have a benzoyl (or substituted benzoyl) group linked via a pyrrolyl group and an
25 C₂alkylene group (-CH=CH- or -CH₂CH₂-) to a carbamic acid group. The compounds apparently showed HDAC inhibitory activity in the micromolar range.

Sulfonamides

Oxamflatin, also known as (2E)-5-[3-[(phenylsulfonyl)amino] phenyl]-pent-2-en-4-ynohydroxamic acid, shown below, has been reported to have *in vitro* antiproliferative activity against various mouse and human tumour cell lines, and *in vivo* antitumour activity against B16 melanoma (see, e.g., Sonoda et al., 1996; Kim et al., 1999).

Oxamflatin



- 10 Ohtani et al., 1993, describe a number of hydroxamic acid compounds which are claimed to be inhibitors of ras transformation. Many of the compounds are hydroxmic acid compounds which have a sulfonamide group, and which employ an acid leader which is: a phenylene-ortho-alkylene (e.g., I-10); phenylene-meta-alkylene (e.g., I-24); phenylene-para-alkylene (e.g., I-12); or naphthylen-1,2-diyl
- 15 (e.g., I-20). However, in every case, the sulfonamide group is -SO₂NR-, as opposed to -NRSO₂-. Also, in every case, the terminal aryl group is linked directly to the -SO₂NR- sulfonamide group, without an intervening aryl leader. Ohtani et al., 1996, describe similar compounds.
- 20 Richon et al., 2001, describe various branched compounds which apparently inhibit histone deacetylase. See the table at pages 96-101 therein. Some of the compounds are carbamic acid compounds having a carbamic acid group (-CONHOH) linked to a branch point, from which two aryl groups are appended. A few linear carbamic acid compounds are also described, including a single
- 25 -SO₂NH- sulfonamide carbamic acid with a -(CH₂)₅- acid leader (compound 671).

Delorme et al., 2001, describe various carbamic acid compounds, including compounds having, inter alia, a sulfonamide group. Of the 108 compounds in the table at pages 114-123 therein, 88 are carbamic acids (-CONHOH), and the

remainder are terminal amides, -CONHR. Of the 88 carbamic acid compounds, 54 have a sulfonamide linkage.

Of the 54 sulfonamide carbamic acids, 51 are indicated to have a -SO₂NR- sulfonamide group, and 3 (compounds 98, 161, and 162) are indicated to have a -NRSO₂- sulfonamide group.

All of the 54 sulfonamide carbamic acids employ a phenylene-alkylene acid leader group (analogous to Q² herein). Of the 54 compounds, 52 employ a phenylene-para-alkylene group, and only 2 (compounds 41 and 26) employ a phenylene-meta-alkylene group (-Ph-CH₂- and -Ph-(CH₂)₄-, respectively). Compounds 41 and 26 both have a -SO₂NR- sulfonamide group, as opposed to a -NRSO₂- sulfonamide group; the former has a benzothiophenyl group, and the latter has a phenyl group.

All but one of the 54 sulfonamide carbamic acids have an aryl group linked directly to the sulfonamide; compound 100 has a benzyl group (Ph-CH₂-) linked a -SO₂NR- sulfonamide group linked to phenylene-para-ethylene.

SUMMARY OF THE INVENTION

One aspect of the invention pertains to active carbamic acid compounds, as described herein, which inhibit HDAC activity.

Another aspect of the invention pertains to active compounds, as described herein, which treat a proliferative condition, such as cancer or psoriasis.

Another aspect of the invention pertains to active compounds, as described herein, which treat conditions which are known to be mediated by HDAC, or which are known to be treated by HDAC inhibitors (such as, e.g., trichostatin A).

Another aspect of the present invention pertains to a composition comprising a compound as described herein and a pharmaceutically acceptable carrier.

Another aspect of the present invention pertains to methods of inhibiting HDAC in a cell, comprising contacting said cell with an effective amount of an active compound, as described herein.

5

Another aspect of the present invention pertains to methods of inhibiting cell proliferation, comprising contacting a cell with an effective amount of an active compound, as described herein, whether *in vitro* or *in vivo*.

10

Another aspect of the present invention pertains to methods of treating a proliferative condition in a patient comprising administering to said patient a therapeutically-effective amount of an active compound, as described herein. In one preferred embodiment, the proliferative condition is cancer. In one preferred embodiment, the proliferative condition is psoriasis.

15

Another aspect of the present invention pertains to methods of treating a condition in a patient which is known to be mediated by HDAC, or which is known to be treated by HDAC inhibitors (such as, e.g., trichostatin A), comprising administering to said patient a therapeutically-effective amount of an active

20

compound, as described herein.

Another aspect of the present invention pertains to an active compound, as described herein, for use in a method of treatment of the human or animal body.

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Another aspect of the present invention pertains to use of an active compound, as described herein, for the manufacture of a medicament for use in the treatment of a proliferative condition. In one preferred embodiment, the proliferative condition is cancer. In one preferred embodiment, the proliferative condition is psoriasis.

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Another aspect of the present invention pertains to use of an active compound for the manufacture of a medicament, for example, for the treatment of conditions which are known to be mediated by HDAC, or which are known to be treated by HDAC inhibitors (such as, e.g., trichostatin A), as discussed herein.

Another aspect of the present invention pertains to a kit comprising (a) the active compound, preferably provided as a pharmaceutical composition and in a suitable container and/or with suitable packaging; and (b) instructions for use, for example, written instructions on how to administer the active compound.

Another aspect of the present invention pertains to compounds obtainable by a method of synthesis as described herein, or a method comprising a method of synthesis as described herein.

Another aspect of the present invention pertains to compounds obtained by a method of synthesis as described herein, or a method comprising a method of synthesis as described herein.

Another aspect of the present invention pertains to novel intermediates, as described herein, which are suitable for use in the methods of synthesis described herein.

Another aspect of the present invention pertains to the use of such novel intermediates, as described herein, in the methods of synthesis described herein.

As will be appreciated by one of skill in the art, features and preferred embodiments of one aspect of the invention will also pertain to other aspects of the invention.

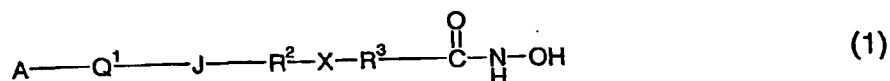
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DETAILED DESCRIPTION OF THE INVENTION

Compounds

In one embodiment, the present invention pertains to carbamic acid compounds of the formula:

- 15 -

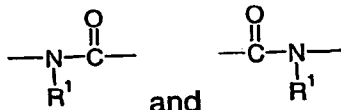


wherein:

A is an aryl group;

Q^1 is a covalent bond or an aryl leader group;

J is an amide linkage selected from:



5

R^1 is an amido substituent;

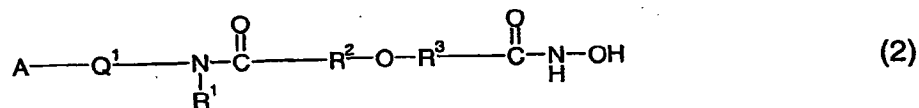
X is an ether heteroatom, and is -O- or -S-; and,

R^2 and R^3 are each independently an ether group;

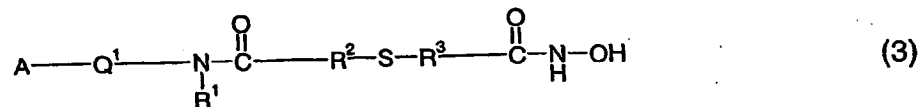
and pharmaceutically acceptable salts, solvates, amides, esters, ethers,
10 chemically protected forms, and prodrugs thereof.

In preferred embodiments, the carbamic acid group, $-\text{C}(=\text{O})\text{NHOH}$, is unmodified
(e.g., is not an ester).

15 In one preferred embodiment, J is $-\text{NR}^1\text{CO}-$ and X is -O-, and the compounds have the following formula:



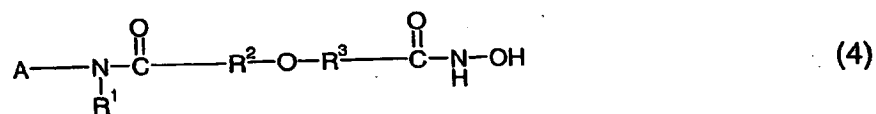
In one preferred embodiment, J is $-\text{NR}^1\text{CO}-$ and X is -S-, and the compounds have the following formula:



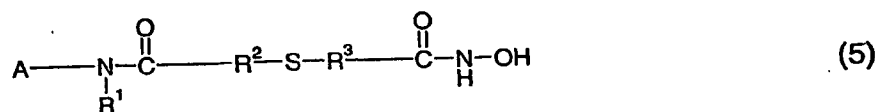
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In one preferred embodiment, Q^1 is a covalent bond, J is $-\text{NR}^1\text{CO}-$ and X is -O-, and the compounds have the following formula:

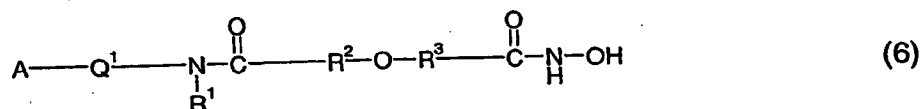
- 16 -



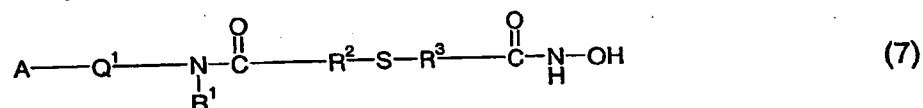
In one preferred embodiment, Q¹ is a covalent bond, J is -NR¹CO- and X is -S-, and the compounds have the following formula:



- 5 In one preferred embodiment, Q¹ is an aryl leader group, J is -NR¹CO- and X is -O-, and the compounds have the following formula:

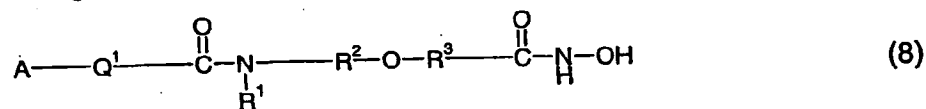


In one preferred embodiment, Q¹ is an aryl leader group, J is -NR¹CO- and X is -S-, and the compounds have the following formula:

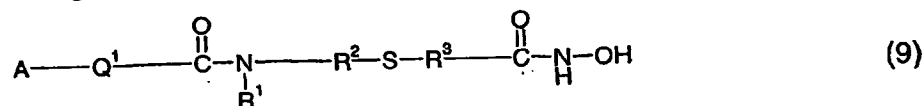


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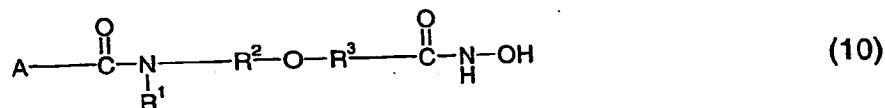
In one preferred embodiment, J is -CONR¹- and X is -O-, and the compounds have the following formula:



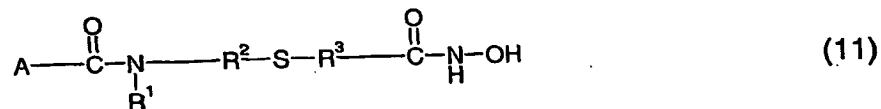
- 15 In one preferred embodiment, J is -CONR¹- and X is -S-, and the compounds have the following formula:



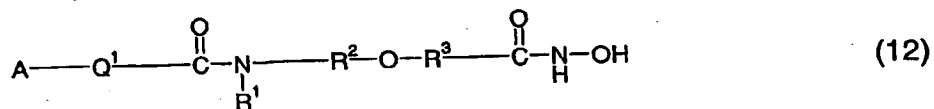
In one preferred embodiment, Q¹ is a covalent bond, J is -CONR¹- and X is -O-, and the compounds have the following formula:



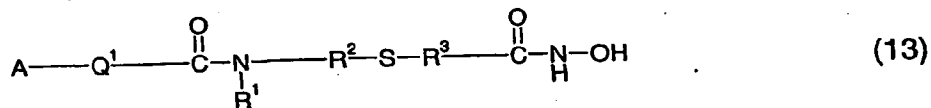
In one preferred embodiment, Q^1 is a covalent bond, J is $-\text{CONR}^1-$ and X is $-\text{S}-$, and the compounds have the following formula:



- 5 In one preferred embodiment, Q^1 is an aryl leader group, J is $-\text{CONR}^1-$ and X is $-\text{O}-$, and the compounds have the following formula:



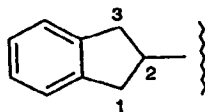
In one preferred embodiment, Q^1 is an aryl leader group, J is $-\text{CONR}^1-$ and X is $-\text{S}-$, and the compounds have the following formula:



10

In one embodiment, where Q^1 is an aryl leader, the aryl group, A, is linked to Q^1 via a covalent single bond.

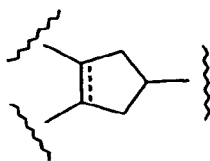
- 15 In one embodiment, where Q^1 is a cyclic aryl leader, the aryl group, A, may be fused to Q^1 and so the moiety A-Q^1- forms a fused polycyclic structure. For example, the moiety 2,3-dihydro-1H-indene-2-yl, derived from indan (2,3-dihydro-1H-indene), is considered to be a phenyl group (A) fused to a C_5 cycloalkyl group (Q^1):



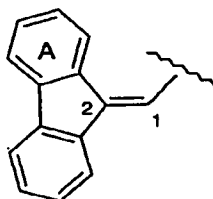
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In such cases, the tridentate aryl leader, Q^1 , may be denoted as:

- 18 -

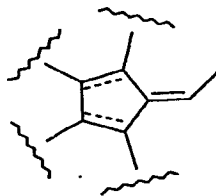


In a similar example, the moiety 9H-fluorene-9-yl, derived from fluorene, is considered to be two phenyl groups (either of which is A), fused to a C₅cycloalkyl group, which forms part of Q¹:



5

In such cases, the pentadentate aryl leader, Q¹, may be denoted as:



10 The Aryl Group, A

The aryl group, A, is a C₅₋₂₀aryl group, and is optionally substituted.

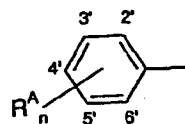
15 In one preferred embodiment, A is a C₅₋₂₀heteroaryl group, and is optionally substituted. In one preferred embodiment, A is a monocyclic C₅₋₂₀heteroaryl group, and is optionally substituted. In one preferred embodiment, A is a monocyclic C₅₋₆heteroaryl group, and is optionally substituted.

20 In one preferred embodiment, A is a C₅₋₂₀carboaryl group, and is optionally substituted. In one preferred embodiment, A is a monocyclic C₅₋₂₀carboaryl group, and is optionally substituted. In one preferred embodiment, A is a monocyclic C₅₋₆carboaryl group, and is optionally substituted. In one preferred embodiment, A is a phenyl group, and is optionally substituted.

In one preferred embodiment, A is a C₅₋₂₀aryl group derived from one of the following: benzene, pyridine, furan, indole, pyrrole, imidazole, naphthalene, quinoline, benzimidazole, benzothiofuran, fluorene, acridine, and carbazole.

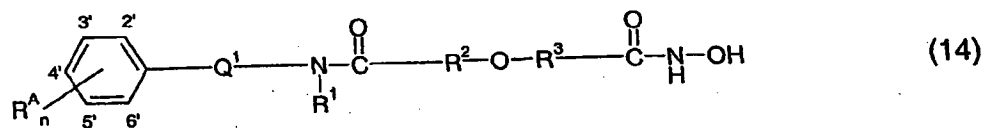
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In one preferred embodiment, A is an optionally substituted phenyl group of the formula:

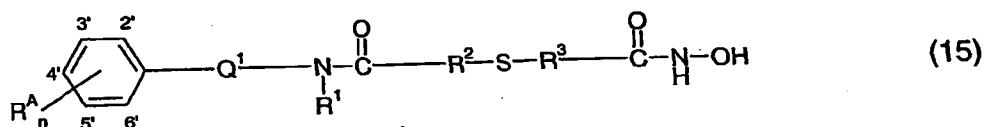


10 wherein n is an integer from 0 to 5, and each R^A is independently a substituent as defined herein.

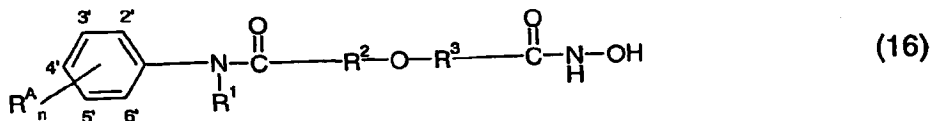
In one preferred embodiment, A is an optionally substituted phenyl group, J is -NR¹CO- and X is -O-, and the compounds have the following formula:



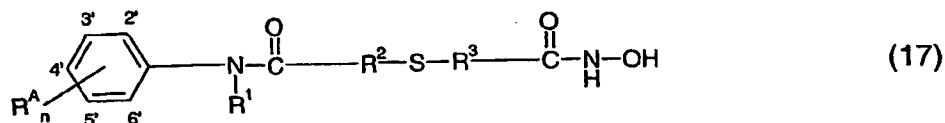
15 In one preferred embodiment, A is an optionally substituted phenyl group, J is -NR¹CO- and X is -S-, and the compounds have the following formula:



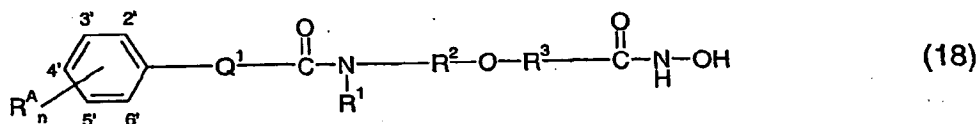
20 In one preferred embodiment, A is an optionally substituted phenyl group, Q¹ is a covalent bond, J is -NR¹CO- and X is -O-, and the compounds have the following formula:



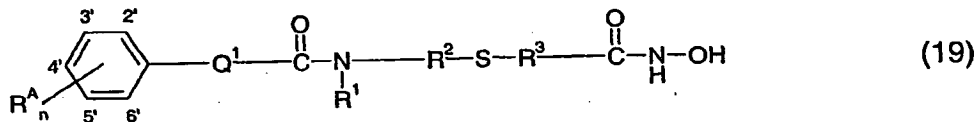
In one preferred embodiment, A is an optionally substituted phenyl group, Q¹ is a covalent bond, J is -NR¹CO- and X is -S-, and the compounds have the following formula:



- 5 In one preferred embodiment, A is an optionally substituted phenyl group, J is -CONR¹- and X is -O-, and the compounds have the following formula:

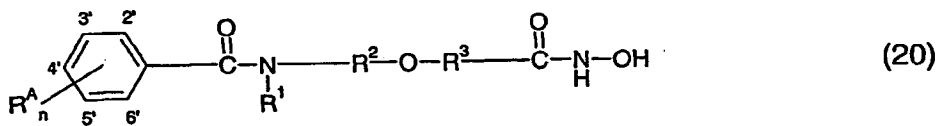


In one preferred embodiment, A is an optionally substituted phenyl group, J is -CONR¹- and X is -S-, and the compounds have the following formula:

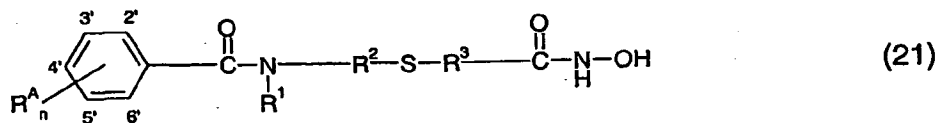


10

In one preferred embodiment, A is an optionally substituted phenyl group, Q¹ is a covalent bond, J is -CONR¹- and X is -O-, and the compounds have the following formula:



- 15 In one preferred embodiment, A is an optionally substituted phenyl group, Q¹ is a covalent bond, J is -CONR¹- and X is -S-, and the compounds have the following formula:



In one preferred embodiment, n is an integer from 0 to 5. In one preferred embodiment, n is an integer from 0 to 4. In one preferred embodiment, n is an integer from 0 to 3. In one preferred embodiment, n is an integer from 0 to 2. In one preferred embodiment, n is 0 or 1.

- 5 In one preferred embodiment, n is an integer from 1 to 5.
In one preferred embodiment, n is an integer from 1 to 4.
In one preferred embodiment, n is an integer from 1 to 3.
In one preferred embodiment, n is 1 or 2.
In one preferred embodiment, n is 5.

- 10 In one preferred embodiment, n is 4.
In one preferred embodiment, n is 3.
In one preferred embodiment, n is 2.
In one preferred embodiment, n is 1.
In one preferred embodiment, n is 0.

15

If the phenyl group has less than the full complement of ring substituents, R^A , they may be arranged in any combination. For example, if n is 1, R^A may be in the 2'-, 3'-, 4'-, 5'-, or 6'-position. Similarly, if n is 2, the two R^A groups may be in, for example, the 2',3'-, 2',4'-, 2',5'-, 2',6'-, 3',4'-, or 3',5'-positions. If n is 3, the three

20 R^A groups may be in, for example, the 2',3',4'-, 2',3',5'-, 2',3',6'-, or 3',4',5'-positions.

In one preferred embodiment, n is 1, and the R^A group is in the 4'-position.

- 25 In one preferred embodiment, n is 2, and one R^A group is in the 4'-position, and the other R^A group is in the 2'-position.

In one preferred embodiment, n is 2, and one R^A group is in the 4'-position, and the other R^A group is in the 3'-position.

30

Each aryl substituent, R^A , is a substituent as defined herein.

Examples of preferred aryl substituents, R^A , include, but are not limited to, the following: fluoro, chloro, bromo, iodo, methyl, ethyl, isopropyl, t-butyl, cyano, trifluoromethyl, hydroxy, methoxy, ethoxy, isopropoxy, trifluoromethoxy, phenoxy, methylthio, trifluoromethylthio, hydroxymethyl, amino, dimethylamino, diethylamino, morpholino, amido (unsubstituted, i.e., $-\text{CONH}_2$), acetamido, acetyl, nitro, sulfonamido (unsubstituted, i.e., $-\text{SO}_2\text{NH}_2$), and phenyl.

In one preferred embodiment, A is a substituted phenyl group selected from:

- para-(fluoro)phenyl; ortho-(fluoro)phenyl; meta-(fluoro)phenyl;
- 10 para-(chloro)phenyl; ortho-(chloro)phenyl; meta-(chloro)phenyl;
- para-(bromo)phenyl; ortho-(bromo)phenyl; meta-(bromo)phenyl;
- para-(iodo)phenyl; ortho-(iodo)phenyl; meta-(iodo)phenyl;
- para-(methyl)phenyl; ortho-(methyl)phenyl; meta-(methyl)phenyl;
- para-(ethyl)phenyl; ortho-(ethyl)phenyl; meta-(ethyl)phenyl;
- 15 para-(isopropyl)phenyl; ortho-(isopropyl)phenyl; meta-(isopropyl)phenyl;
- para-(t-butyl)phenyl; ortho-(t-butyl)phenyl; meta-(t-butyl)phenyl;
- para-(cyano)phenyl; ortho-(cyano)phenyl; meta-(cyano)phenyl;
- para-(trifluoromethyl)phenyl; ortho-(trifluoromethyl)phenyl; meta-
- (trifluoromethyl)phenyl;
- 20 para-(hydroxy)phenyl; ortho-(hydroxy)phenyl; meta-(hydroxy)phenyl;
- para-(methoxy)phenyl; ortho-(methoxy)phenyl; meta-(methoxy)phenyl;
- para-(ethoxy)phenyl; ortho-(ethoxy)phenyl; meta-(ethoxy)phenyl;
- para-(isopropoxy)phenyl; ortho-(isopropoxy)phenyl;
- meta-(isopropoxy)phenyl;
- 25 para-(trifluoromethoxy)phenyl; ortho-(trifluoromethoxy)phenyl;
- meta-(trifluoromethoxy)phenyl;
- para-(phenoxy)phenyl; ortho-(phenoxy)phenyl; meta-(phenoxy)phenyl;
- para-(methylthio)phenyl; ortho-(methylthio)phenyl; meta-(methylthio)phenyl;
- para-(trifluoromethylthio)phenyl; ortho-(trifluoromethylthio)phenyl;
- 30 meta-(trifluoromethylthio)phenyl;
- para-(hydroxymethyl)phenyl; ortho-(hydroxymethyl)phenyl;
- meta-(hydroxymethyl)phenyl;
- para-(amino)phenyl; ortho-(amino)phenyl; meta-(amino)phenyl;

para-(dimethylamino)phenyl; ortho-(dimethylamino)phenyl;
meta-(dimethylamino)phenyl;
para-(diethylamino)phenyl; ortho-(diethylamino)phenyl;
meta-(diethylamino)phenyl;
5 para-(morpholino)phenyl; ortho-(morpholino)phenyl;
meta-(morpholino)phenyl;
para-(amido)phenyl; ortho-(amido)phenyl; meta-(amido)phenyl;
para-(acetamido)phenyl; ortho-(acetamido)phenyl;
meta-(acetamido)phenyl;
10 para-(acetyl)phenyl; ortho-(acetyl)phenyl; meta-(acetyl)phenyl;
para-(nitro)phenyl; ortho-(nitro)phenyl; meta-(nitro)phenyl;
para-(sulfonamido)phenyl; ortho-(sulfonamido)phenyl;
meta-(sulfonamido)phenyl; and,
para-(phenyl)phenyl; ortho-(phenyl)phenyl; meta-(phenyl)phenyl.

15

In one preferred embodiment, A is a substituted phenyl group selected from:

para-(fluoro)phenyl;
para-(chloro)phenyl;
para-(bromo)phenyl;
20 para-(iodo)phenyl;
para-(methyl)phenyl;
para-(ethyl)phenyl;
para-(isopropyl)phenyl;
para-(t-butyl)phenyl;
25 para-(cyano)phenyl;
para-(trifluoromethyl)phenyl;
para-(hydroxy)phenyl;
para-(methoxy)phenyl;
para-(ethoxy)phenyl;
30 para-(isopropoxy)phenyl;
para-(trifluoromethoxy)phenyl;
para-(phenoxy)phenyl;
para-(methylthio)phenyl;

- para-(trifluoromethylthio)phenyl;
para-(hydroxymethyl)phenyl;
para-(amino)phenyl;
para-(dimethylamino)phenyl;
5 para-(diethylamino)phenyl;
para-(morpholino)phenyl;
para-(amido)phenyl;
para-(acetamido)phenyl;
para-(acetyl)phenyl;
10 para-(nitro)phenyl;
para-(sulfonamido)phenyl; and,
para-(phenyl)phenyl.

In one preferred embodiment, A is a substituted phenyl group selected from:

- 15 ortho,para-di(methoxy)phenyl;
ortho,para-di(halo)phenyl;
ortho,para-di(fluoro)phenyl;
ortho-(methoxy),para-(methyl)phenyl;
20 ortho-(methoxy),para-(trifluoromethyl)phenyl;
ortho-(trifluoromethyl),para-(halo)phenyl;

ortho,meta-di(trifluoromethyl)phenyl;
ortho-(halo),meta-(trifluoromethyl)phenyl;
25
meta,para-di(halo)phenyl;
meta,para-di(hydroxy)phenyl;
meta,para-di(methyl)phenyl;
meta,para-di(methoxy)phenyl;
30 meta-(halo),para-(nitro)phenyl;

3',5'-di(trifluoromethyl)phenyl;
3'-(trifluoromethyl),5'-(methoxy)phenyl;

- 25 -

3'-(trifluoromethyl),5'-(halo)phenyl;

2'-(halo),5'-(methyl)phenyl;

5 2',6'-di(methyl)phenyl;
 2',6'-di(halo)phenyl;
 2',6'-di(isopropyl)phenyl;

10 2',4',6'-tri(halo)phenyl;
 3',4',5'-tri(halo)phenyl;
 3',4',5'-tri(methoxy)phenyl;
 2',5'-di(halo)-4'-(hydroxy)phenyl; and
 3'-(trifluoromethyl),5',6'-di(halo)phenyl.

15 The Aryl Leader Group, Q¹

As mentioned above, in some embodiments, Q¹ is a covalent bond or an aryl leader group; in some embodiments, Q¹ is a covalent bond; in some embodiments, Q¹ is an aryl leader group.

20 In one preferred embodiment, Q¹ is a covalent bond.

In one preferred embodiment, Q¹ is a C₁₋₇alkylene group and is optionally substituted.

25 In one preferred embodiment, Q¹ is a covalent bond or a C₁₋₇alkylene group and is optionally substituted.

30 In one preferred embodiment, Q¹ is a covalent bond or a saturated C₁₋₇alkylene group. In one preferred embodiment, Q¹ is a saturated C₁₋₇alkylene group.

In one preferred embodiment, Q¹ is a covalent bond or a partially unsaturated C₁₋₇alkylene group. In one preferred embodiment, Q¹ is a partially unsaturated C₁₋₇alkylene group.

- 5 In one preferred embodiment, Q¹ is a covalent bond or an aliphatic C₁₋₇alkylene group. In one preferred embodiment, Q¹ is an aliphatic C₁₋₇alkylene group.

In one preferred embodiment, Q¹ is a covalent bond or a linear C₁₋₇alkylene group. In one preferred embodiment, Q¹ is a linear C₁₋₇alkylene group.

10

In one preferred embodiment, Q¹ is a covalent bond or a branched C₁₋₇alkylene group. In one preferred embodiment, Q¹ is a branched C₁₋₇alkylene group.

- 15 In one preferred embodiment, Q¹ is a covalent bond or an alicyclic C₁₋₇alkylene group. In one preferred embodiment, Q¹ is an alicyclic C₁₋₇alkylene group.

In one preferred embodiment, Q¹ is a covalent bond or a saturated aliphatic C₁₋₇alkylene group. In one preferred embodiment, Q¹ is a saturated aliphatic C₁₋₇alkylene group.

20

In one preferred embodiment, Q¹ is a covalent bond or a saturated linear C₁₋₇alkylene group. In one preferred embodiment, Q¹ is a saturated linear C₁₋₇alkylene group.

- 25 In one preferred embodiment, Q¹ is a covalent bond or a saturated branched C₁₋₇alkylene group. In one preferred embodiment, Q¹ is a saturated branched C₁₋₇alkylene group.

- 30 In one preferred embodiment, Q¹ is a covalent bond or a saturated alicyclic C₁₋₇alkylene group. In one preferred embodiment, Q¹ is a saturated alicyclic C₁₋₇alkylene group.

In one preferred embodiment, Q¹ is a covalent bond or a partially unsaturated aliphatic C₁₋₇alkylene group. In one preferred embodiment, Q¹ is a partially unsaturated aliphatic C₁₋₇alkylene group.

- 5 In one preferred embodiment, Q¹ is a covalent bond or a partially unsaturated linear C₁₋₇alkylene group. In one preferred embodiment, Q¹ is a partially unsaturated linear C₁₋₇alkylene group.

- 10 In one preferred embodiment, Q¹ is a covalent bond or a partially unsaturated branched C₁₋₇alkylene group. In one preferred embodiment, Q¹ is a partially unsaturated branched C₁₋₇alkylene group.

- 15 In one preferred embodiment, Q¹ is a covalent bond or a partially unsaturated alicyclic C₁₋₇alkylene group. In one preferred embodiment, Q¹ is a partially unsaturated alicyclic C₁₋₇alkylene group.

- Note that, as discussed below in the context of isomers, where unsaturation permits isomers, e.g., cis- and trans, E- and Z-, etc., and combinations thereof, a reference to one isomer is to be considered a reference to all such isomers,
20 unless otherwise specified.

The Aryl Leader Group, Q¹: Substituents

- In one embodiment, Q¹ is unsubstituted.
25 In one embodiment, Q¹ is optionally substituted.
In one embodiment, Q¹ is substituted.

Examples of substituents on Q¹ include, but are not limited to, those described under the heading "Substituents" below.

- 30 In one preferred embodiment, substituents on Q¹, if present, are independently selected from: halo, hydroxy, ether (e.g., C₁₋₇alkoxy), C₅₋₂₀aryl, acyl, amido, and oxo.

In one preferred embodiment, substituents on Q^1 , if present, are independently selected from -F, -Cl, -Br, -I, -OH, -OMe, -OEt, -OPr, -Ph, and =O.

- 5 In one preferred embodiment, substituents on Q^1 , if present, are -OH or -Ph.

In one preferred embodiment, substituents on Q^1 , if present, are -Ph.

- For example, in one embodiment, Q^1 is unsubstituted ethylene, and is $-\text{CH}_2\text{CH}_2-$;
10 in one embodiment, Q^1 is oxo ($=\text{O}$) substituted ethylene, and is $-\text{C}(=\text{O})\text{CH}_2-$; in one embodiment, Q^1 is hydroxy ($-\text{OH}$) substituted ethylene, and is $-\text{CH}(\text{OH})\text{CH}_2-$; in one embodiment, Q^1 is phenyl ($-\text{Ph}$) substituted ethylene, and is $-\text{CH}_2\text{CH}(\text{Ph})-$.

The Aryl Leader Group, Q^1 : Certain Embodiments

15

Note that, for embodiments excluding, e.g., a covalent bond, certain backbone lengths, etc., it is to be understood that the corresponding species listed below are similarly excluded from the respective embodiments discussed below.

- 20 In one preferred embodiment, Q^1 is selected from the following:

a covalent bond;

$-(\text{CH}_2)_n-$ where n is an integer from 1 to 7;

25

$-\text{CH}(\text{CH}_3)-$;

$-\text{CH}(\text{CH}_3)\text{CH}_2-$ and $-\text{CH}_2\text{CH}(\text{CH}_3)-$;

$-\text{CH}(\text{CH}_3)\text{CH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}(\text{CH}_3)\text{CH}_2-$, and $-\text{CH}_2\text{CH}_2\text{CH}(\text{CH}_3)-$;

$-\text{CH}(\text{CH}_3)\text{CH}_2\text{CH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}(\text{CH}_3)\text{CH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}_2\text{CH}(\text{CH}_3)\text{CH}_2-$, and

- 30 $-\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}(\text{CH}_3)-$;

$-\text{CH}(\text{CH}_3)\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}(\text{CH}_3)\text{CH}_2\text{CH}_2\text{CH}_2-$,

$-\text{CH}_2\text{CH}_2\text{CH}(\text{CH}_3)\text{CH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}(\text{CH}_3)\text{CH}_2-$, and

$-\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}(\text{CH}_3)-$;

- $-\text{CH}(\text{CH}_2\text{CH}_3)-;$
 $-\text{CH}(\text{CH}_2\text{CH}_3)\text{CH}_2-$ and $-\text{CH}_2\text{CH}(\text{CH}_2\text{CH}_3)-;$
 $-\text{CH}(\text{CH}_2\text{CH}_3)\text{CH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}(\text{CH}_2\text{CH}_3)\text{CH}_2-$, and $-\text{CH}_2\text{CH}_2\text{CH}(\text{CH}_2\text{CH}_3)-;$
5 $-\text{CH}(\text{CH}_2\text{CH}_3)\text{CH}_2\text{CH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}(\text{CH}_2\text{CH}_3)\text{CH}_2\text{CH}_2-$,
 $-\text{CH}_2\text{CH}_2\text{CH}(\text{CH}_2\text{CH}_3)\text{CH}_2-$, and $-\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}(\text{CH}_2\text{CH}_3)-;$
 $-\text{CH}(\text{CH}_2\text{CH}_3)\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}(\text{CH}_2\text{CH}_3)\text{CH}_2\text{CH}_2\text{CH}_2-$,
 $-\text{CH}_2\text{CH}_2\text{CH}(\text{CH}_2\text{CH}_3)\text{CH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}(\text{CH}_2\text{CH}_3)\text{CH}_2-$, and
 $-\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}(\text{CH}_2\text{CH}_3)-;$
10
 $-\text{CH}=\text{CH}-;$
 $-\text{CH}=\text{CHCH}_2-$ and $-\text{CH}_2\text{CH}=\text{CH}-;$
 $-\text{CH}=\text{CHCH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}=\text{CHCH}_2-$, and $-\text{CH}_2\text{CH}_2\text{CH}=\text{CH}-;$
 $-\text{CH}=\text{CHCH}_2\text{CH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}=\text{CHCH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}_2\text{CH}=\text{CHCH}_2-$, and
15 $-\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}=\text{CH}-;$
 $-\text{CH}=\text{CHCH}_2\text{CH}_2\text{CH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}=\text{CHCH}_2\text{CH}_2\text{CH}_2-$,
 $-\text{CH}_2\text{CH}_2\text{CH}=\text{CHCH}_2\text{CH}_2-$, $-\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}=\text{CHCH}_2-$, and
 $-\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_2\text{CH}=\text{CH}-;$
20
 $-\text{C}(\text{CH}_3)=\text{CH}-$ and $-\text{CH}=\text{C}(\text{CH}_3)-;$
 $-\text{C}(\text{CH}_3)=\text{CHCH}_2-$, $-\text{CH}=\text{C}(\text{CH}_3)\text{CH}_2-$, and $-\text{CH}=\text{CHCH}(\text{CH}_3)-;$
 $-\text{CH}(\text{CH}_3)\text{CH}=\text{CH}-$, $-\text{CH}_2\text{C}(\text{CH}_3)=\text{CH}-$, and $-\text{CH}_2\text{CH}=\text{C}(\text{CH}_3)-;$
25
 $-\text{CH}=\text{CHCH}=\text{CH}-;$
 $-\text{CH}=\text{CHCH}=\text{CHCH}_2-$, $-\text{CH}_2\text{CH}=\text{CHCH}=\text{CH}-$, and $-\text{CH}=\text{CHCH}_2\text{CH}=\text{CH}-;$
 $-\text{CH}=\text{CHCH}=\text{CHCH}_2\text{CH}_2-$, $-\text{CH}=\text{CHCH}_2\text{CH}=\text{CHCH}_2-$,
 $-\text{CH}=\text{CHCH}_2\text{CH}_2\text{CH}=\text{CH}-$, $-\text{CH}_2\text{CH}=\text{CHCH}=\text{CHCH}_2-$, $-\text{CH}_2\text{CH}=\text{CHCH}_2\text{CH}=\text{CH}-$,
and $-\text{CH}_2\text{CH}_2\text{CH}=\text{CHCH}=\text{CH}-;$
30
 $-\text{C}(\text{CH}_3)=\text{CHCH}=\text{CH}-$, $-\text{CH}=\text{C}(\text{CH}_3)\text{CH}=\text{CH}-$, $-\text{CH}=\text{CHC}(\text{CH}_3)=\text{CH}-$, and
 $-\text{CH}=\text{CHCH}=\text{C}(\text{CH}_3)-;$
 $-\text{C}\equiv\text{C}-;$

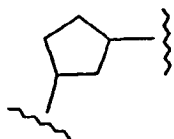
- C \equiv CCH₂-, -CH₂C \equiv C-, -C \equiv CCH(CH₃)-, and -CH(CH₃)C \equiv C-;
 -C \equiv CCH₂CH₂-, -CH₂C \equiv CCH₂-, and -CH₂CH₂C \equiv C-;
 -C \equiv CCH(CH₃)CH₂- and -C \equiv CCH₂CH(CH₃)-;
 -CH(CH₃)C \equiv CCH₂- and -CH₂C \equiv CCH(CH₃)-;
 5 -CH(CH₃)CH₂C \equiv C- and -CH₂CH(CH₃)C \equiv C-;
 -C \equiv CCH=CH-, -CH=CHC \equiv C-, and -C \equiv CC \equiv C-;
 -C \equiv CCH₂CH₂CH₂- and -CH₂CH₂CH₂C \equiv C-;
 -C \equiv CCH₂CH₂CH₂CH₂- and -CH₂CH₂CH₂CH₂C \equiv C-;
 -C \equiv CCH=CHCH=CH-, -CH=CHC \equiv C-CH=CH-, and -CH=CHCH=CHC \equiv C-;
 10 -C(CH₃)=CHC \equiv C-, -CH=C(CH₃)C \equiv C-, -C \equiv CC(CH₃)=CH-, and
 -C \equiv CCH=C(CH₃)-;

- cyclopentylene and cyclopentenylene; and,
 15 cyclohexylene, cyclohexenylene, and cyclohexadienylene.

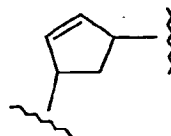
In one preferred embodiment, Q¹ is selected from:

- a covalent bond;
 -CH₂-, -(CH₂)₂-, -(CH₂)₃-, -(CH₂)₄-, -(CH₂)₅-, and -(CH₂)₆-;
 20 -CH(CH₃)CH₂CH₂CH₂CH₂-, -CH₂CH(CH₃)CH₂CH₂CH₂-,
 -CH₂CH₂CH₂CH(CH₃)CH₂-, and -CH₂CH₂CH₂CH₂CH(CH₃)-;
 -CH=CH-;
 -CH=CH-CH=CH-;
 -CH=CHCH₂CH₂CH₂- and -CH₂CH₂CH₂CH=CH-;
 25 -CH=CHCH₂CH₂CH₂CH₂- and -CH₂CH₂CH₂CH₂CH=CH-;
 -C(CH₃)=CHCH=CH-, -CH=C(CH₃)CH=CH-, -CH=CHC(CH₃)=CH-, and
 -CH=CHCH=C(CH₃)-;

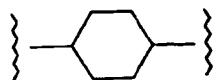
(cyclopent-1,3-ylene)



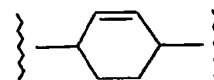
(4-cyclopenten-1,3-ylene)



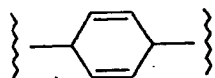
(cyclohex-1,4-ylylene)



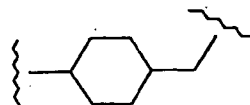
(2-cyclohexen-1,4-ylylene)



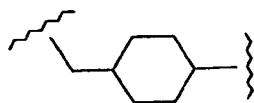
(2,5-cyclohexadien-1,4-ylylene)



(cyclohex-1,4-ylylene-methylene)



(methylene-cyclohex-1,4-ylylene)



In one preferred embodiment, Q¹ is selected from:

a covalent bond;

-CH₂-, -(CH₂)₂-, -(CH₂)₃-, -(CH₂)₄-, -(CH₂)₅-,

5

-CH=CH-;

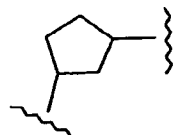
-CH=CH-CH=CH-;

-C(CH₃)=CHCH=CH-, -CH=C(CH₃)CH=CH-, -CH=CHC(CH₃)=CH-, and

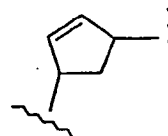
-CH=CHCH=C(CH₃)-;

-CH=CHCH₂CH₂CH₂- and -CH₂CH₂CH₂CH=CH-; and,

(cyclopent-1,3-ylylene)



(4-cyclopenten-1,3-ylylene)



10

In one preferred embodiment, Q¹ is selected from: a covalent bond, -CH₂-, -CH₂CH₂-, -CH₂CH₂CH₂-, -CH=CH-, and -CH=CH-CH=CH-.

In one preferred embodiment, Q¹ is selected from:

15

a covalent bond, -CH₂-, -CH=CH-, and -CH=CH-CH=CH-.

In one preferred embodiment, Q¹ is selected from:

-CH₂-, -CH=CH-, and -CH=CH-CH=CH-.

The Amido Substituent, R¹

- The amido substituent, R¹, is hydrogen, C₁₋₇alkyl (including, e.g., C₅₋₂₀aryl-C₁₋₇alkyl), C₃₋₂₀heterocyclyl, or C₅₋₂₀aryl.

Note that R¹ is a monodentate species. It is not intended that R¹ be additionally linked to A, Q¹, and/or Q², thereby forming a cyclic group.

- 10 In one preferred embodiment, R¹ is hydrogen, C₁₋₇alkyl, or C₅₋₂₀aryl.
In one preferred embodiment, R¹ is hydrogen or C₁₋₇alkyl.

In one preferred embodiment, R¹ is hydrogen, saturated C₁₋₇alkyl, or C₅₋₂₀aryl.

In one preferred embodiment, R¹ is hydrogen or saturated C₁₋₇alkyl.

15

In one preferred embodiment, R¹ is hydrogen, saturated aliphatic C₁₋₇alkyl, or C₅₋₂₀aryl.

In one preferred embodiment, R¹ is hydrogen or saturated aliphatic C₁₋₇alkyl.

- 20 In one preferred embodiment, R¹ is -H, -Me, -Et, -nPr, -iPr, -nBu, -sBu, -tBu, -Ph, or -Bn.

In one preferred embodiment, R¹ is -H, -Me, -Et, -nPr, -iPr, -nBu, -sBu, or -tBu.

In one preferred embodiment, R¹ is -H, -Me, -Et, -Ph, or -Bn.

- 25 In one preferred embodiment, R¹ is -H, -Me, or -Et.

In one preferred embodiment, R¹ is -H.

The Ether Groups, R² and R³

30

Each of the ether groups, R² and R³, is independently C₁₋₇alkylene; C₅₋₂₀arylene; C₅₋₂₀arylene-C₁₋₇alkylene; or C₁₋₇alkylene-C₅₋₂₀arylene; and is optionally substituted.

In one preferred embodiment, each of the ether groups, R^2 and R^3 , is independently a C_{1-7} alkylene group, and is optionally substituted.

- 5 In one embodiment, each of R^2 and R^3 is unsubstituted.
In one embodiment, each of R^2 and R^3 is optionally substituted.
In one embodiment, each of R^2 and R^3 is substituted.

The Ether Groups, R^2 and R^3 : Alkylene

10

In one preferred embodiment, R^2 and/or R^3 is a saturated C_{1-7} alkylene group.

In one preferred embodiment, R^2 and/or R^3 is a partially unsaturated C_{1-7} alkylene group.

15

In one preferred embodiment, R^2 and/or R^3 is an aliphatic C_{1-7} alkylene group.

In one preferred embodiment, R^2 and/or R^3 is a linear C_{1-7} alkylene group.

- 20 In one preferred embodiment, R^2 and/or R^3 is a branched C_{1-7} alkylene group.

In one preferred embodiment, R^2 and/or R^3 is an alicyclic C_{1-7} alkylene group.

- 25 In one preferred embodiment, R^2 and/or R^3 is a saturated aliphatic C_{1-7} alkylene group.

In one preferred embodiment, R^2 and/or R^3 is a saturated linear C_{1-7} alkylene group.

- 30 In one preferred embodiment, R^2 and/or R^3 is a saturated branched C_{1-7} alkylene group.

In one preferred embodiment, R^2 and/or R^3 is a saturated alicyclic C_{1-7} alkylene group.

5 In one preferred embodiment, R^2 and/or R^3 is a partially unsaturated aliphatic C_{1-7} alkylene group.

In one preferred embodiment, R^2 and/or R^3 is a partially unsaturated linear C_{1-7} alkylene group.

10 In one preferred embodiment, R^2 and/or R^3 is a partially unsaturated branched C_{1-7} alkylene group.

In one preferred embodiment, R^2 and/or R^3 is a partially unsaturated alicyclic C_{1-7} alkylene group.

15

In one preferred embodiment, R^2 and/or R^3 is selected from:

$-(CH_2)_n-$ where n is an integer from 1 to 7;

20

$-CH(CH_3)-$;

$-CH(CH_3)CH_2-$ and $-CH_2CH(CH_3)-$;

$-CH(CH_3)CH_2CH_2-$, $-CH_2CH(CH_3)CH_2-$, and $-CH_2CH_2CH(CH_3)-$;

$-CH(CH_3)CH_2CH_2CH_2-$, $-CH_2CH(CH_3)CH_2CH_2-$, $-CH_2CH_2CH(CH_3)CH_2-$, and

$-CH_2CH_2CH_2CH(CH_3)-$;

25

$-CH(CH_3)CH_2CH_2CH_2CH_2-$, $-CH_2CH(CH_3)CH_2CH_2CH_2-$,

$-CH_2CH_2CH(CH_3)CH_2CH_2-$, $-CH_2CH_2CH_2CH(CH_3)CH_2-$, and

$-CH_2CH_2CH_2CH_2CH(CH_3)-$;

$-CH(CH_2CH_3)-$;

30

$-CH(CH_2CH_3)CH_2-$ and $-CH_2CH(CH_2CH_3)-$;

$-CH(CH_2CH_3)CH_2CH_2-$, $-CH_2CH(CH_2CH_3)CH_2-$, and $-CH_2CH_2CH(CH_2CH_3)-$;

$-CH(CH_2CH_3)CH_2CH_2CH_2-$, $-CH_2CH(CH_2CH_3)CH_2CH_2-$,

$-CH_2CH_2CH(CH_2CH_3)CH_2-$, and $-CH_2CH_2CH_2CH(CH_2CH_3)-$;

-CH(CH₂CH₃)CH₂CH₂CH₂CH₂-, -CH₂CH(CH₂CH₃)CH₂CH₂CH₂-,
 -CH₂CH₂CH(CH₂CH₃)CH₂CH₂-, -CH₂CH₂CH₂CH(CH₂CH₃)CH₂-, and
 -CH₂CH₂CH₂CH₂CH(CH₂CH₃)-

5 -CH=CH-;
 -CH=CHCH₂- and -CH₂CH=CH-;
 -CH=CHCH₂CH₂-, -CH₂CH=CHCH₂-, and -CH₂CH₂CH=CH-;
 -CH=CHCH₂CH₂CH₂-, -CH₂CH=CHCH₂CH₂-, -CH₂CH₂CH=CHCH₂-, and
 10 -CH₂CH₂CH₂CH=CH-;

 -CH=CHCH₂CH₂CH₂CH₂-, -CH₂CH=CHCH₂CH₂CH₂-,
 -CH₂CH₂CH=CHCH₂CH₂-, -CH₂CH₂CH₂CH=CHCH₂-, and
 -CH₂CH₂CH₂CH₂CH=CH-;

 -C(CH₃)=CH- and -CH=C(CH₃)-;
 15 -C(CH₃)=CHCH₂-, -CH=C(CH₃)CH₂-, and -CH=CHCH(CH₃)-;
 -CH(CH₃)CH=CH-, -CH₂C(CH₃)=CH-, and -CH₂CH=C(CH₃)-

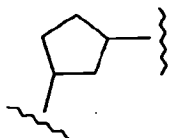
 -CH=CHCH=CH-;
 -CH=CHCH=CHCH₂-, -CH₂CH=CHCH=CH-, and -CH=CHCH₂CH=CH-;
 20 -CH=CHCH=CHCH₂CH₂-, -CH=CHCH₂CH=CHCH₂-, and
 -CH=CHCH₂CH₂CH=CH-, -CH₂CH=CHCH=CHCH₂-, -CH₂CH=CHCH₂CH=CH-,
 and -CH₂CH₂CH=CHCH=CH-;

 cyclopentylene and cyclopentenylene; and,
 25 cyclohexylene, cyclohexenylene, and cyclohexadienylene.

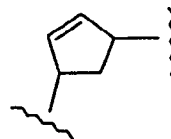
In one preferred embodiment, R² and/or R³ is selected from:

 -CH₂-, -(CH₂)₂-, -(CH₂)₃-, -(CH₂)₄-, -(CH₂)₅-, and -(CH₂)₆;
 -CH(CH₃)CH₂CH₂CH₂CH₂-, -CH₂CH(CH₃)CH₂CH₂CH₂-,
 30 -CH₂CH₂CH₂CH(CH₃)CH₂-, and -CH₂CH₂CH₂CH₂CH(CH₃)-;
 -CH=CHCH₂CH₂CH₂- and -CH₂CH₂CH₂CH=CH-;
 -CH=CHCH₂CH₂CH₂CH₂- and -CH₂CH₂CH₂CH₂CH=CH-;

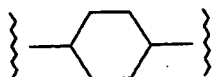
(cyclopent-1,3-ylylene)



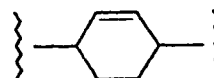
(4-cyclopenten-1,3-ylylene)



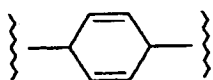
(cyclohex-1,4-ylylene)



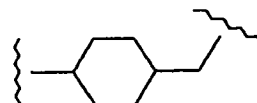
(2-cyclohexen-1,4-ylylene)



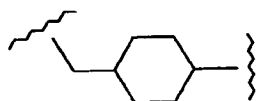
(2,5-cyclohexadien-1,4-ylylene)



(cyclohex-1,4-ylylene-methylene)



(methylene-cyclohex-1,4-ylylene)



In one preferred embodiment, each of R^2 and R^3 is a saturated C_{1-7} alkylene group.

- 5 In one preferred embodiment, each of R^2 and R^3 is selected from $-(CH_2)_n-$, wherein n is an integer from 1 to 5.

In one preferred embodiment, the group R^2-X-R^3 is selected from the following:

- 10 $-CH_2-O-CH_2-$ and $-CH_2-S-CH_2-$;
 $-CH_2-O-CH_2CH_2-$ and $-CH_2-S-CH_2CH_2-$;
 $-CH_2CH_2-O-CH_2-$ and $-CH_2CH_2-S-CH_2-$;
 $-CH_2-O-CH_2CH_2CH_2-$ and $-CH_2-S-CH_2CH_2CH_2-$;
 $-CH_2CH_2-O-CH_2CH_2-$ and $-CH_2CH_2-S-CH_2CH_2-$;
 $-CH_2CH_2CH_2-O-CH_2-$ and $-CH_2CH_2CH_2-S-CH_2-$;
15 $-CH_2-O-CH_2CH_2CH_2CH_2-$ and $-CH_2-S-CH_2CH_2CH_2CH_2-$;
 $-CH_2CH_2-O-CH_2CH_2CH_2-$ and $-CH_2CH_2-S-CH_2CH_2CH_2-$;
 $-CH_2CH_2CH_2-O-CH_2CH_2-$ and $-CH_2CH_2CH_2-S-CH_2CH_2-$;
 $-CH_2CH_2CH_2CH_2-O-CH_2-$ and $-CH_2CH_2CH_2CH_2-S-CH_2-$;

- 5 -CH₂-O-CH₂CH₂CH₂CH₂CH₂- and -CH₂-S-CH₂CH₂CH₂CH₂CH₂-;
 -CH₂CH₂-O-CH₂CH₂CH₂CH₂- and -CH₂CH₂-S-CH₂CH₂CH₂CH₂-;
 -CH₂CH₂CH₂-O-CH₂CH₂CH₂- and -CH₂CH₂CH₂-S-CH₂CH₂CH₂-;
 -CH₂CH₂CH₂CH₂-O-CH₂CH₂- and -CH₂CH₂CH₂CH₂-S-CH₂CH₂-;
 -CH₂CH₂CH₂CH₂CH₂-O-CH₂- and -CH₂CH₂CH₂CH₂CH₂-S-CH₂-;

In one preferred embodiment, the group R²-X-R³ is selected from the following:

-CH₂-O-CH₂- and -CH₂-S-CH₂-.

- 10 In one preferred embodiment, the group R²-X-R³ is selected from the following:

-CH₂-O-CH₂CH₂- and -CH₂-S-CH₂CH₂-;

-CH₂CH₂-O-CH₂- and -CH₂CH₂-S-CH₂-.

In one preferred embodiment, the group R²-X-R³ is selected from the following:

- 15 -CH₂-O-CH₂CH₂CH₂- and -CH₂-S-CH₂CH₂CH₂-;

-CH₂CH₂-O-CH₂CH₂- and -CH₂CH₂-S-CH₂CH₂-;

-CH₂CH₂CH₂-O-CH₂- and -CH₂CH₂CH₂-S-CH₂-.

In one preferred embodiment, the group R²-X-R³ is selected from the following:

- 20 -CH₂-O-CH₂CH₂CH₂CH₂- and -CH₂-S-CH₂CH₂CH₂CH₂-;

-CH₂CH₂-O-CH₂CH₂CH₂- and -CH₂CH₂-S-CH₂CH₂CH₂-;

-CH₂CH₂CH₂-O-CH₂CH₂- and -CH₂CH₂CH₂-S-CH₂CH₂-;

-CH₂CH₂CH₂CH₂-O-CH₂- and -CH₂CH₂CH₂CH₂-S-CH₂-.

- 25 In one preferred embodiment, the group R²-X-R³ is selected from the following:

-CH₂-O-CH₂CH₂CH₂CH₂CH₂- and -CH₂-S-CH₂CH₂CH₂CH₂CH₂-;

-CH₂CH₂-O-CH₂CH₂CH₂CH₂- and -CH₂CH₂-S-CH₂CH₂CH₂CH₂-;

-CH₂CH₂CH₂-O-CH₂CH₂CH₂- and -CH₂CH₂CH₂-S-CH₂CH₂CH₂-;

-CH₂CH₂CH₂CH₂-O-CH₂CH₂- and -CH₂CH₂CH₂CH₂-S-CH₂CH₂-;

- 30 -CH₂CH₂CH₂CH₂CH₂-O-CH₂- and -CH₂CH₂CH₂CH₂CH₂-S-CH₂-.

The Ether Groups, R² and R³: Arylene

In one preferred embodiment, R^2 and/or R^3 is C_{5-20} arylene, and is optionally substituted.

- 5 In one preferred embodiment, R^2 and/or R^3 is C_{5-20} arylene. In one preferred embodiment, R^2 and/or R^3 is C_{5-6} arylene. In one preferred embodiment, R^2 and/or R^3 is phenylene.

The Ether Groups, R^2 and R^3 : Alkylene-Arylene and Arylene-Alkylene

- 10 In one preferred embodiment, R^2 and/or R^3 is C_{5-20} arylene- C_{1-7} alkylene or C_{1-7} alkylene- C_{5-20} arylene, and is optionally substituted.

In one preferred embodiment, R^2 and/or R^3 is C_{5-6} arylene- C_{1-7} alkylene or C_{1-7} alkylene- C_{5-6} arylene, and is optionally substituted.

15

In one preferred embodiment, R^2 and/or R^3 is C_{1-7} alkylene- C_{5-20} arylene.

In one preferred embodiment, R^2 and/or R^3 is C_{1-7} alkylene- C_{5-6} arylene.

In one preferred embodiment, R^2 and/or R^3 is C_{5-20} arylene- C_{1-7} alkylene.

- 20 In one preferred embodiment, R^2 and/or R^3 is C_{5-6} arylene- C_{1-7} alkylene.

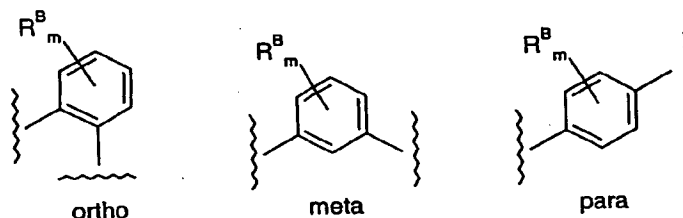
In one preferred embodiment, R^2 and/or R^3 is C_{1-7} alkylene-phenylene. In one preferred embodiment, R^2 and/or R^3 is methylene-phenylene, ethylene-phenylene, propylene-phenylene, and ethenylene-phenylene (also known as vinylene-

- 25 phenylene).

In one preferred embodiment, R^2 and/or R^3 is phenylene- C_{1-7} alkylene. In one preferred embodiment, R^2 and/or R^3 is phenylene-methylene, phenylene-ethylene, phenylene-propylene, or phenylene-ethenylene (also known as phenylene-

- 30 vinylene).

In the above alkylene-phenylene and phenylene-alkylene groups, the phenylene linkage may be ortho, meta, or para, and the phenylene group is optionally substituted with from 1 to 4 aryl substituents, R^B :



5

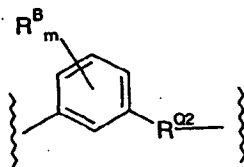
In one preferred embodiment, the phenylene linkage is meta or para. In one preferred embodiment, the phenylene linkage is para. In one preferred embodiment, the phenylene linkage is meta.

- 10 In one preferred embodiment, m is an integer from 0 to 4.
In one preferred embodiment, m is an integer from 0 to 3.
In one preferred embodiment, m is an integer from 0 to 2.
In one preferred embodiment, m is 0 or 1.
In one preferred embodiment, m is an integer from 1 to 4.
- 15 In one preferred embodiment, m is an integer from 1 to 3.
In one preferred embodiment, m is 1 or 2.
In one preferred embodiment, m is 4.
In one preferred embodiment, m is 3.
In one preferred embodiment, m is 2.
- 20 In one preferred embodiment, m is 1.
In one preferred embodiment, m is 0.

Each aryl substituent, R^B , is a substituent as defined herein.

- 25 Examples of preferred aryl substituents, R^B , include, but are not limited to, the following: fluoro, chloro, methyl, ethyl, isopropyl, *t*-butyl, trifluoromethyl, hydroxy, methoxy, ethoxy, isopropoxy, methylthio, amino, dimethylamino, diethylamino, morpholino, acetamido, nitro, and phenyl.

In one preferred embodiment, the phenylene linkage is meta, and R^2 and/or R^3 has the following formula, wherein R^{Q2} is C_{1-7} alkylene and is optionally substituted (referred to herein as "phenylene-meta- C_{1-7} alkylene"):



5

In one preferred embodiment, R^{Q2} is a saturated C_{1-7} alkylene group.

In one preferred embodiment, R^{Q2} is a partially unsaturated C_{1-7} alkylene group.

10 In one preferred embodiment, R^{Q2} is an aliphatic C_{1-7} alkylene group.

In one preferred embodiment, R^{Q2} is a linear C_{1-7} alkylene group.

In one preferred embodiment, R^{Q2} is a branched C_{1-7} alkylene group.

15

In one preferred embodiment, R^{Q2} is an alicyclic C_{1-7} alkylene group.

In one preferred embodiment, R^{Q2} is a saturated aliphatic C_{1-7} alkylene group.

20 In one preferred embodiment, R^{Q2} is a saturated linear C_{1-7} alkylene group.

In one preferred embodiment, R^{Q2} is a saturated branched C_{1-7} alkylene group.

In one preferred embodiment, R^{Q2} is a saturated alicyclic C_{1-7} alkylene group.

25

In one preferred embodiment, R^{Q2} is a partially unsaturated aliphatic C_{1-7} alkylene group.

30 In one preferred embodiment, R^{Q2} is a partially unsaturated linear C_{1-7} alkylene group.

In one preferred embodiment, R^{Q2} is a partially unsaturated branched C_{1-7} alkylene group.

- 5 In one preferred embodiment, R^{Q2} is a partially unsaturated alicyclic C_{1-7} alkylene group.

In one preferred embodiment, R^{Q2} is selected from:

- 10 $-(CH_2)_n-$ where n is an integer from 1 to 7;
 $-CH(CH_3)-$;
 $-CH(CH_3)CH_2-$ and $-CH_2CH(CH_3)-$;
 $-CH(CH_3)CH_2CH_2-$, $-CH_2CH(CH_3)CH_2-$, and $-CH_2CH_2CH(CH_3)-$;
 $-CH(CH_3)CH_2CH_2CH_2-$, $-CH_2CH(CH_3)CH_2CH_2-$, $-CH_2CH_2CH(CH_3)CH_2-$, and
- 15 $-CH_2CH_2CH_2CH(CH_3)-$;
 $-CH(CH_3)CH_2CH_2CH_2CH_2-$, $-CH_2CH(CH_3)CH_2CH_2CH_2-$,
 $-CH_2CH_2CH(CH_3)CH_2CH_2-$, $-CH_2CH_2CH_2CH(CH_3)CH_2-$, and
 $-CH_2CH_2CH_2CH_2CH(CH_3)-$;
- 20 $-CH(CH_2CH_3)-$;
 $-CH(CH_2CH_3)CH_2-$ and $-CH_2CH(CH_2CH_3)-$;
 $-CH(CH_2CH_3)CH_2CH_2-$, $-CH_2CH(CH_2CH_3)CH_2-$, and $-CH_2CH_2CH(CH_2CH_3)-$;
 $-CH(CH_2CH_3)CH_2CH_2CH_2-$, $-CH_2CH(CH_2CH_3)CH_2CH_2-$,
 $-CH_2CH_2CH(CH_2CH_3)CH_2-$, and $-CH_2CH_2CH_2CH(CH_2CH_3)-$;
- 25 $-CH(CH_2CH_3)CH_2CH_2CH_2CH_2-$, $-CH_2CH(CH_2CH_3)CH_2CH_2CH_2-$,
 $-CH_2CH_2CH(CH_2CH_3)CH_2CH_2-$, $-CH_2CH_2CH_2CH(CH_2CH_3)CH_2-$, and
 $-CH_2CH_2CH_2CH_2CH(CH_2CH_3)-$;
- 30 $-CH=CH-$;
 $-CH=CHCH_2-$ and $-CH_2CH=CH-$;
 $-CH=CHCH_2CH_2-$, $-CH_2CH=CHCH_2-$, and $-CH_2CH_2CH=CH-$;
 $-CH=CHCH_2CH_2CH_2-$, $-CH_2CH=CHCH_2CH_2-$, $-CH_2CH_2CH=CHCH_2-$, and
 $-CH_2CH_2CH_2CH=CH-$;

-CH=CHCH₂CH₂CH₂CH₂-, -CH₂CH=CHCH₂CH₂CH₂-,
 -CH₂CH₂CH=CHCH₂CH₂-, -CH₂CH₂CH₂CH=CHCH₂-, and
 -CH₂CH₂CH₂CH₂CH=CH-;

5 -C(CH₃)=CH- and -CH=C(CH₃)-;
 -C(CH₃)=CHCH₂-, -CH=C(CH₃)CH₂-, and -CH=CHCH(CH₃)-;
 -CH(CH₃)CH=CH-, -CH₂C(CH₃)=CH-, and -CH₂CH=C(CH₃)-;

10 -CH=CHCH=CH-;
 -CH=CHCH=CHCH₂-, -CH₂CH=CHCH=CH-, and -CH=CHCH₂CH=CH-;
 -CH=CHCH=CHCH₂CH₂-, -CH=CHCH₂CH=CHCH₂-, and
 -CH=CHCH₂CH₂CH=CH-, -CH₂CH=CHCH=CHCH₂-, -CH₂CH=CHCH₂CH=CH-,
 and -CH₂CH₂CH=CHCH=CH-;

15 -C(CH₃)=CHCH=CH-, -CH=C(CH₃)CH=CH-, -CH=CHC(CH₃)=CH-, and
 -CH=CHCH=C(CH₃)-;

 cyclopentylene and cyclopentenylene; and,
 cyclohexylene, cyclohexenylene, and cyclohexadienylene.

20

In one preferred embodiment, R^{Q2} is selected from:

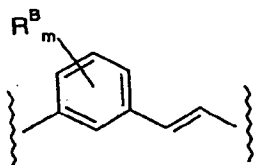
 -CH₂-, -(CH₂)₂-, -(CH₂)₃-, -(CH₂)₄-, -(CH₂)₅-, and -(CH₂)₆-;
 -CH=CH-, -CH=CH-CH=CH-;

25 In one preferred embodiment, R^{Q2} is cis or trans -CH=CH-.

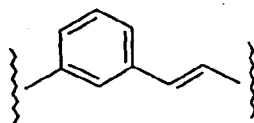
 In one preferred embodiment, R^{Q2} is cis -CH=CH-.

 In one preferred embodiment, R^{Q2} is trans -CH=CH-.

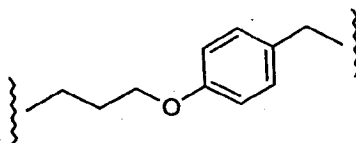
In one preferred embodiment, R^{Q2} is $-\text{CH}=\text{CH}-$, and R^2 and/or R^3 is (referred to herein as "phenylene-meta-trans-ethylene"):



- 5 In one preferred embodiment, m is 0, and R^2 and/or R^3 is (referred to herein as "unsubstituted phenylene-meta-trans-ethylene"):

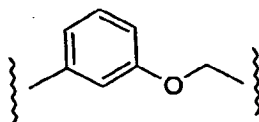


In one preferred embodiment, $-\text{R}^2-\text{X}-\text{R}^3-$ is:



10

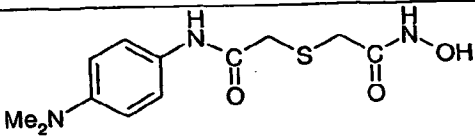
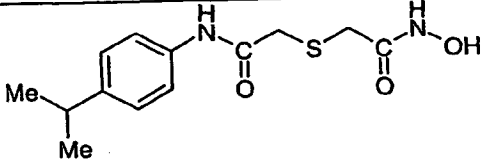
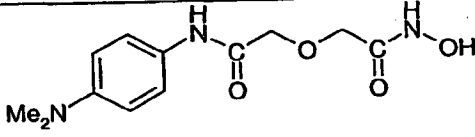
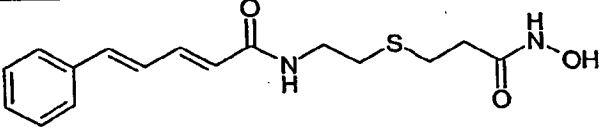
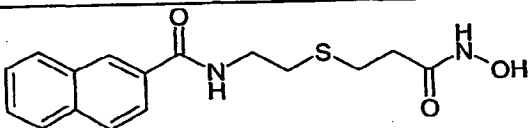
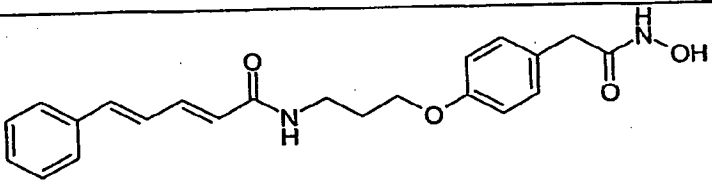
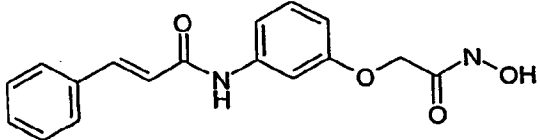
In one preferred embodiment, $-\text{R}^2-\text{X}-\text{R}^3-$ is:



15 Examples of Specific Embodiments

Some individual embodiments of the present invention include the following compounds.

1		PX083805
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2		PX089278
3		PX089279
4		PX089281
5		PX117439
6		PX117440
7		PX117443
8		PX117708

Chemical Terms

The term "carbo," "carbyl," "hydrocarbo," and "hydrocarbyl," as used herein, pertain to compounds and/or groups which have only carbon and hydrogen atoms.

The term "hetero," as used herein, pertains to compounds and/or groups which have at least one heteroatom, for example, multivalent heteroatoms (which are also suitable as ring heteroatoms) such as boron, silicon, nitrogen, phosphorus,

oxygen, and sulfur, and monovalent heteroatoms, such as fluorine, chlorine, bromine, and iodine.

The term "saturated," as used herein, pertains to compounds and/or groups which
5 do not have any carbon-carbon double bonds or carbon-carbon triple bonds.

The term "unsaturated," as used herein, pertains to compounds and/or groups which have at least one carbon-carbon double bond or carbon-carbon triple bond.

10 The term "aliphatic," as used herein, pertains to compounds and/or groups which are linear or branched, but not cyclic (also known as "acyclic" or "open-chain" groups).

The term "cyclic," as used herein, pertains to compounds and/or groups which
15 have one ring, or two or more rings (e.g., spiro, fused, bridged).

The term "ring," as used herein, pertains to a closed ring of from 3 to 10 covalently linked atoms, more preferably 3 to 8 covalently linked atoms.

20 The term "aromatic ring," as used herein, pertains to a closed ring of from 3 to 10 covalently linked atoms, more preferably 5 to 8 covalently linked atoms, which ring is aromatic.

The term "heterocyclic ring," as used herein, pertains to a closed ring of from 3 to
25 10 covalently linked atoms, more preferably 3 to 8 covalently linked atoms, wherein at least one of the ring atoms is a multivalent ring heteroatom, for example, nitrogen, phosphorus, silicon, oxygen, and sulfur, though more commonly nitrogen, oxygen, and sulfur.

30 The term "alicyclic," as used herein, pertains to compounds and/or groups which have one ring, or two or more rings (e.g., spiro, fused, bridged), wherein said ring(s) are not aromatic.

The term "aromatic," as used herein, pertains to compounds and/or groups which have one ring, or two or more rings (e.g., fused), wherein at least one of said ring(s) is aromatic.

- 5 The term "heterocyclic," as used herein, pertains to cyclic compounds and/or groups which have one heterocyclic ring, or two or more heterocyclic rings (e.g., spiro, fused, bridged), wherein said ring(s) may be alicyclic or aromatic.

- 10 The term "heteroaromatic," as used herein, pertains to cyclic compounds and/or groups which have one heterocyclic ring, or two or more heterocyclic rings (e.g., fused), wherein said ring(s) is aromatic.

Substituents

- 15 The phrase "optionally substituted," as used herein, pertains to a parent group which may be unsubstituted or which may be substituted.

- 20 Unless otherwise specified, the term "substituted," as used herein, pertains to a parent group which bears one or more substituents. The term "substituent" is used herein in the conventional sense and refers to a chemical moiety which is covalently attached to, appended to, or if appropriate, fused to, a parent group. A wide variety of substituents are well known, and methods for their formation and introduction into a variety of parent groups are also well known.

- 25 In one preferred embodiment, the substituent(s), often referred to herein as R, are independently selected from: halo; hydroxy; ether (e.g., C₁₋₇alkoxy); formyl; acyl (e.g., C₁₋₇alkylacyl, C₅₋₂₀arylacyl); acylhalide; carboxy; ester; acyloxy; amido; acylamido; thioamido; tetrazolyl; amino; nitro; nitroso; azido; cyano; isocyano; cyanato; isocyanato; thiocyano; isothiocyano; sulfhydryl; thioether (e.g., C₁₋₇alkylthio); sulfonic acid; sulfonate; sulfone; sulfonyloxy; sulfinyloxy; sulfamino; 30 sulfonamino; sulfinamino; sulfamyl; sulfonamido; C₁₋₇alkyl (including, e.g., C₁₋₇haloalkyl, C₁₋₇hydroxyalkyl, C₁₋₇carboxyalkyl, C₁₋₇aminoalkyl, C₅₋₂₀aryl-

C₁₋₇alkyl); C₃₋₂₀heterocyclyl; or C₅₋₂₀aryl (including, e.g., C₅₋₂₀carboaryl, C₅₋₂₀heteroaryl, C₁₋₇alkyl-C₅₋₂₀aryl and C₅₋₂₀haloaryl)).

In one preferred embodiment, the substituent(s), often referred to herein as R, are
5 independently selected from:

- F, -Cl, -Br, and -I;
- OH;
- OMe, -OEt, -O(tBu), and -OCH₂Ph;
- SH;
- 10 -SMe, -SEt, -S(tBu), and -SCH₂Ph;
- C(=O)H;
- C(=O)Me, -C(=O)Et, -C(=O)(tBu), and -C(=O)Ph;
- C(=O)OH;
- C(=O)OMe, -C(=O)OEt, and -C(=O)O(tBu);
- 15 -C(=O)NH₂, -C(=O)NHMe, -C(=O)NMe₂, and -C(=O)NHEt;
- NHC(=O)Me, -NHC(=O)Et, -NHC(=O)Ph, succinimidyl, and maleimidyl;
- NH₂, -NHMe, -NH₂Et, -NH(iPr), -NH(nPr), -NMe₂, -NEt₂, -N(iPr)₂, -N(nPr)₂,
-N(nBu)₂, and -N(tBu)₂;
- CN;
- 20 -NO₂;
- Me, -Et, -nPr, -iPr, -nBu, -tBu;
- CF₃, -CHF₂, -CH₂F, -CCl₃, -CBr₃, -CH₂CH₂F, -CH₂CHF₂, and -CH₂CF₃;
- OCF₃, -OCHF₂, -OCH₂F, -OCCl₃, -OCBr₃, -OCH₂CH₂F, -OCH₂CHF₂, and
-OCH₂CF₃;
- 25 -CH₂OH, -CH₂CH₂OH, and -CH(OH)CH₂OH;
- CH₂NH₂, -CH₂CH₂NH₂, and -CH₂CH₂NMe₂; and,
optionally substituted phenyl.

In one preferred embodiment, the substituent(s), often referred to herein as R, are
30 independently selected from: -F, -Cl, -Br, -I, -OH, -OMe, -OEt, -SH, -SMe, -SEt,
-C(=O)Me, -C(=O)OH, -C(=O)OMe, -CONH₂, -CONHMe, -NH₂, -NMe₂, -NEt₂,
-N(nPr)₂, -N(iPr)₂, -CN, -NO₂, -Me, -Et, -CF₃, -OCF₃, -CH₂OH, -CH₂CH₂OH,
-CH₂NH₂, -CH₂CH₂NH₂, and -Ph.

In one preferred embodiment, the substituent(s), often referred to herein as R, are independently selected from: hydroxy; ether (e.g., C₁₋₇alkoxy); ester; amido; amino; and, C₁₋₇alkyl (including, e.g., C₁₋₇haloalkyl, C₁₋₇hydroxyalkyl,

- 5 C₁₋₇carboxyalkyl, C₁₋₇aminoalkyl, C₅₋₂₀aryl-C₁₋₇alkyl).

In one preferred embodiment, the substituent(s), often referred to herein as R, are independently selected from:

- OH;
- 10 -OMe, -OEt, -O(tBu), and -OCH₂Ph;
-C(=O)OMe, -C(=O)OEt, and -C(=O)O(tBu);
-C(=O)NH₂, -C(=O)NHMe, -C(=O)NMe₂, and -C(=O)NHET;
-NH₂, -NHMe, -NHET, -NH(iPr), -NH(nPr), -NMe₂, -NEt₂, -N(iPr)₂, -N(nPr)₂,
-N(nBu)₂, and -N(tBu)₂;
- 15 -Me, -Et, -nPr, -iPr, -nBu, -tBu;
-CF₃, -CHF₂, -CH₂F, -CCl₃, -CBr₃, -CH₂CH₂F, -CH₂CHF₂, and -CH₂CF₃;
-CH₂OH, -CH₂CH₂OH, and -CH(OH)CH₂OH; and,
-CH₂NH₂, -CH₂CH₂NH₂, and -CH₂CH₂NMe₂.
- 20 The substituents are described in more detail below.

- C₁₋₇alkyl: The term "C₁₋₇alkyl," as used herein, pertains to a monovalent moiety obtained by removing a hydrogen atom from a C₁₋₇hydrocarbon compound having from 1 to 7 carbon atoms, which may be aliphatic or alicyclic, or a combination thereof, and which may be saturated, partially unsaturated, or fully unsaturated.
- 25

Examples of (unsubstituted) saturated linear C₁₋₇alkyl groups include, but are not limited to, methyl, ethyl, n-propyl, n-butyl, and n-pentyl (amyl).

- 30 Examples of (unsubstituted) saturated branched C₁₋₇alkyl groups include, but are not limited to, iso-propyl, iso-butyl, sec-butyl, tert-butyl, and neo-pentyl.

Examples of saturated alicyclic (also carbocyclic) C₁₋₇alkyl groups (also referred to as "C₃₋₇cycloalkyl" groups) include, but are not limited to, unsubstituted groups such as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, and norbornane, as well as substituted groups (e.g., groups which comprise such groups), such as

5 methylcyclopropyl, dimethylcyclopropyl, methylcyclobutyl, dimethylcyclobutyl, methylcyclopentyl, dimethylcyclopentyl, methylcyclohexyl, dimethylcyclohexyl, cyclopropylmethyl and cyclohexylmethyl.

Examples of (unsubstituted) unsaturated C₁₋₇alkyl groups which have one or more carbon-carbon double bonds (also referred to as "C₂₋₇alkenyl" groups) include, but

10 are not limited to, ethenyl (vinyl, -CH=CH₂), 2-propenyl (allyl, -CH-CH=CH₂), isopropenyl (-C(CH₃)=CH₂), butenyl, pentenyl, and hexenyl.

Examples of (unsubstituted) unsaturated C₁₋₇alkyl groups which have one or more carbon-carbon triple bonds (also referred to as "C₂₋₇alkynyl" groups) include, but

15 are not limited to, ethynyl (ethinyl) and 2-propynyl (propargyl).

Examples of unsaturated alicyclic (also carbocyclic) C₁₋₇alkyl groups which have one or more carbon-carbon double bonds (also referred to as "C₃₋₇cycloalkenyl" groups) include, but are not limited to, unsubstituted groups such as

20 cyclopropenyl, cyclobutenyl, cyclopentenyl, and cyclohexenyl, as well as substituted groups (e.g., groups which comprise such groups) such as cyclopropenylmethyl and cyclohexenylmethyl.

Additional examples of substituted C₃₋₇cycloalkyl groups include, but are not limited to, those with one or more other rings fused thereto, for example, those derived from: indene (C₉), indan (2,3-dihydro-1H-indene) (C₉), tetraline (1,2,3,4-tetrahydronaphthalene) (C₁₀), adamantane (C₁₀), decalin (decahydronaphthalene) (C₁₂), fluorene (C₁₃), phenalene (C₁₃). For example, 2H-inden-2-yl is a

25 C₅cycloalkyl group with a substituent (phenyl) fused thereto.

30

C₃₋₂₀heterocyclyl: The term "C₃₋₂₀heterocyclyl," as used herein, pertains to a monovalent moiety obtained by removing a hydrogen atom from a ring atom of a

C₃₋₂₀heterocyclic compound, said compound having one ring, or two or more rings (e.g., spiro, fused, bridged), and having from 3 to 20 ring atoms, of which from 1 to 10 are ring heteroatoms, and wherein at least one of said ring(s) is a heterocyclic ring. Preferably, each ring has from 3 to 7 ring atoms, of which from 1 to 4 are ring heteroatoms.

In this context, the prefixes (e.g., C₃₋₂₀, C₃₋₇, C₅₋₆, etc.) denote the number of ring atoms, or range of number of ring atoms, whether carbon atoms or heteroatoms. For example, the term "C₅₋₆heterocyclyl," as used herein, pertains to a heterocyclyl group having 5 or 6 ring atoms. Examples of groups of heterocyclyl groups include C₃₋₂₀heterocyclyl, C₃₋₇heterocyclyl, C₅₋₇heterocyclyl.

Examples of (non-aromatic) monocyclic heterocyclyl groups include, but are not limited to, those derived from:

15

N₁: aziridine (C₃), azetidine (C₄), pyrrolidine (tetrahydropyrrole) (C₅), pyrroline (e.g., 3-pyrroline, 2,5-dihydropyrrole) (C₅), 2H-pyrrole or 3H-pyrrole (isopyrrole, isoazole) (C₅), piperidine (C₆), dihydropyridine (C₆), tetrahydropyridine (C₆), azepine (C₇);

20

O₁: oxirane (C₃), oxetane (C₄), oxolane (tetrahydrofuran) (C₅), oxole (dihydrofuran) (C₅), oxane (tetrahydropyran) (C₆), dihydropyran (C₆), pyran (C₆), oxepin (C₇);

25

S₁: thiirane (C₃), thietane (C₄), thiolane (tetrahydrothiophene) (C₅), thiane (tetrahydrothiopyran) (C₆), thiepane (C₇);

O₂: dioxolane (C₅), dioxane (C₆), and dioxepane (C₇);

30

O₃: trioxane (C₆);

N₂: imidazolidine (C₅), pyrazolidine (diazolidine) (C₅), imidazoline (C₅), pyrazoline (dihydropyrazole) (C₅), piperazine (C₆);

N₁O₁: tetrahydrooxazole (C₅), dihydrooxazole (C₅), tetrahydroisoxazole (C₅), dihydroisoxazole (C₅), morpholine (C₆), tetrahydrooxazine (C₆), dihydrooxazine (C₆), oxazine (C₆);

5

N₁S₁: thiazoline (C₅), thiazolidine (C₅), thiomorpholine (C₆);

N₂O₁: oxadiazine (C₆);

10 O₁S₁: oxathiole (C₅) and oxathiane (thioxane) (C₆); and,

N₁O₁S₁: oxathiazine (C₆).

15 Examples of substituted (non-aromatic) monocyclic heterocyclyl groups include saccharides, in cyclic form, for example, furanoses (C₅), such as arabinofuranose, lyxofuranose, ribofuranose, and xylofuranse, and pyranoses (C₆), such as allopyranose, altropyranose, glucopyranose, mannopyranose, gulopyranose, idopyranose, galactopyranose, and talopyranose.

20 Examples of heterocyclyl groups which are also heteroaryl groups are described below with aryl groups.

C₅₋₂₀aryl: The term "C₅₋₂₀aryl," as used herein, pertains to a monovalent moiety obtained by removing a hydrogen atom from an aromatic ring atom of a
25 C₅₋₂₀aromatic compound, said compound having one ring, or two or more rings (e.g., fused), and having from 5 to 20 ring atoms, and wherein at least one of said ring(s) is an aromatic ring. Preferably, each ring has from 5 to 7 ring atoms. In this context, the prefixes (e.g., C₃₋₂₀, C₅₋₇, C₅₋₆, etc.) denote the number of ring atoms, or range of number of ring atoms, whether carbon atoms or heteroatoms.
30 For example, the term "C₅₋₆aryl," as used herein, pertains to an aryl group having 5 or 6 ring atoms. Examples of groups of aryl groups include C₃₋₂₀aryl, C₅₋₇aryl, C₅₋₆aryl.

The ring atoms may be all carbon atoms, as in "carboaryl groups" (e.g., C₅₋₂₀carboaryl).

5 Examples of carboaryl groups include, but are not limited to, those derived from benzene (i.e., phenyl) (C₆), naphthalene (C₁₀), azulene (C₁₀), anthracene (C₁₄), phenanthrene (C₁₄), naphthacene (C₁₈), and pyrene (C₁₆).

10 Examples of aryl groups which comprise fused rings, at least one of which is an aromatic ring, include, but are not limited to, groups derived from indene (C₉), isoindene (C₉), and fluorene (C₁₃).

15 Alternatively, the ring atoms may include one or more heteroatoms, including but not limited to oxygen, nitrogen, and sulfur, as in "heteroaryl groups." In this case, the group may conveniently be referred to as a "C₅₋₂₀heteroaryl" group, wherein "C₅₋₂₀" denotes ring atoms, whether carbon atoms or heteroatoms. Preferably, each ring has from 5 to 7 ring atoms, of which from 0 to 4 are ring heteroatoms.

Examples of monocyclic heteroaryl groups include, but are not limited to, those derived from:

20 N₁: pyrrole (azole) (C₅), pyridine (azine) (C₆);

O₁: furan (oxole) (C₅);

S₁: thiophene (thiole) (C₅);

N₁O₁: oxazole (C₅), isoxazole (C₅), isoxazine (C₆);

N₂O₁: oxadiazole (furazan) (C₅);

25 N₃O₁: oxatriazole (C₅);

N₁S₁: thiazole (C₅), isothiazole (C₅);

N₂: imidazole (1,3-diazole) (C₅), pyrazole (1,2-diazole) (C₅), pyridazine (1,2-diazine) (C₆), pyrimidine (1,3-diazine) (C₆) (e.g., cytosine, thymine, uracil), pyrazine (1,4-diazine) (C₆);

30 N₃: triazole (C₅), triazine (C₆); and,

N₄: tetrazole (C₅).

Examples of heterocyclic groups (some of which are also heteroaryl groups) which comprise fused rings, include, but are not limited to:

5 C_9 heterocyclic groups (with 2 fused rings) derived from benzofuran (O_1), isobenzofuran (O_1), indole (N_1), isoindole (N_1), purine (N_4) (e.g., adenine, guanine), benzimidazole (N_2), benzoxazole (N_1O_1), benzisoxazole (N_1O_1), benzodioxole (O_2), benzofurazan (N_2O_1), benzotriazole (N_3), benzothiofuran (S_1), benzothiazole (N_1S_1), benzothiadiazoole (N_2S);

10 C_{10} heterocyclic groups (with 2 fused rings) derived from benzodioxan (O_2), quinoline (N_1), isoquinoline (N_1), benzoxazine (N_1O_1), benzodiazine (N_2), pyridopyridine (N_2), quinoxaline (N_2), quinazoline (N_2);

C_{13} heterocyclic groups (with 3 fused rings) derived from carbazole (N_1), dibenzofuran (O_1), dibenzothiophene (S_1); and,

15 C_{14} heterocyclic groups (with 3 fused rings) derived from acridine (N_1), xanthene (O_1), phenoxathiin (O_1S_1), phenazine (N_2), phenoxazine (N_1O_1), phenothiazine (N_1S_1), thianthrene (S_2), phenanthridine (N_1), phenanthroline (N_2), phenazine (N_2).

20 Heterocyclic groups (including heteroaryl groups) which have a nitrogen ring atom in the form of an -NH- group may be N-substituted, that is, as -NR-. For example, pyrrole may be N-methyl substituted, to give N-methylpyrrole. Examples of N-substitutents include, but are not limited to C_{1-7} alkyl, C_{3-20} heterocyclyl, C_{5-20} aryl, and acyl groups.

25 Heterocyclic groups (including heteroaryl groups) which have a nitrogen ring atom in the form of an -N= group may be substituted in the form of an N-oxide, that is, as -N(\rightarrow O)= (also denoted -N⁺(\rightarrow O⁻)=). For example, quinoline may be substituted to give quinoline N-oxide; pyridine to give pyridine N-oxide; benzofurazan to give benzofurazan N-oxide (also known as benzofuroxan).

30 Cyclic groups may additionally bear one or more oxo (=O) groups on ring carbon atoms. Monocyclic examples of such groups include, but are not limited to, those derived from:

- C₅: cyclopentanone, cyclopentenone, cyclopentadienone;
C₆: cyclohexanone, cyclohexenone, cyclohexadienone;
O₁: furanone (C₅), pyrone (C₆);
N₁: pyrrolidone (pyrrolidinone) (C₅), piperidinone (piperidone) (C₆), piperidinedione
5 (C₆);
N₂: imidazolidone (imidazolidinone) (C₅), pyrazolone (pyrazolinone) (C₅),
piperazinone (C₆), piperazinedione (C₆), pyridazinone (C₆), pyrimidinone (C₆)
(e.g., cytosine), pyrimidinedione (C₆) (e.g., thymine, uracil), barbituric acid (C₆);
N₁S₁: thiazolone (C₅), isothiazolone (C₅);
10 N₁O₁: oxazolinone (C₅).

Polycyclic examples of such groups include, but are not limited to, those derived from:

- C₉: indenedione;
15 N₁: oxindole (C₉);
O₁: benzopyrone (e.g., coumarin, isocoumarin, chromone) (C₁₀);
N₁O₁: benzoxazolinone (C₉), benzoxazolinone (C₁₀);
N₂: quinazolinedione (C₁₀);
N₄: purinone (C₉) (e.g., guanine).

20

Still more examples of cyclic groups which bear one or more oxo (=O) groups on ring carbon atoms include, but are not limited to, those derived from:

- cyclic anhydrides (-C(=O)-O-C(=O)- in a ring), including but not limited to
maleic anhydride (C₅), succinic anhydride (C₅), and glutaric anhydride (C₆);
25 cyclic carbonates (-O-C(=O)-O- in a ring), such as ethylene carbonate (C₅)
and 1,2-propylene carbonate (C₅);
imides (-C(=O)-NR-C(=O)- in a ring), including but not limited to,
succinimide (C₅), maleimide (C₅), phthalimide, and glutarimide (C₆);
lactones (cyclic esters, -O-C(=O)- in a ring), including, but not limited to,
30 β-propiolactone, γ-butyrolactone, δ-valerolactone (2-piperidone), and
ε-caprolactone;

- lactams (cyclic amides, -NR-C(=O)- in a ring), including, but not limited to, β -propiolactam (C₄), γ -butyrolactam (2-pyrrolidone) (C₅), δ -valerolactam (C₆), and ϵ -caprolactam (C₇);
- cyclic carbamates (-O-C(=O)-NR- in a ring), such as 2-oxazolidone (C₅);
- 5 cyclic ureas (-NR-C(=O)-NR- in a ring), such as 2-imidazolidone (C₅) and pyrimidine-2,4-dione (e.g., thymine, uracil) (C₆).

- The above C₁₋₇alkyl, C₃₋₂₀heterocyclyl, and C₅₋₂₀aryl groups, whether alone or part of another substituent, may themselves optionally be substituted with one or more
- 10 groups selected from themselves and the additional substituents listed below.

- Hydrogen: -H. Note that if the substituent at a particular position is hydrogen, it may be convenient to refer to the compound as being "unsubstituted" at that position.

- 15 Halo: -F, -Cl, -Br, and -I.

Hydroxy: -OH.

- 20 Ether: -OR, wherein R is an ether substituent, for example, a C₁₋₇alkyl group (also referred to as a C₁₋₇alkoxy group, discussed below), a C₃₋₂₀heterocyclyl group (also referred to as a C₃₋₂₀heterocyclyloxy group), or a C₅₋₂₀aryl group (also referred to as a C₅₋₂₀aryloxy group), preferably a C₁₋₇alkyl group.

- 25 C₁₋₇alkoxy: -OR, wherein R is a C₁₋₇alkyl group. Examples of C₁₋₇alkoxy groups include, but are not limited to, -OCH₃ (methoxy), -OCH₂CH₃ (ethoxy) and -OC(CH₃)₃ (tert-butoxy).

- Oxo (keto, -one): =O. Examples of cyclic compounds and/or groups having, as a
- 30 substituent, an oxo group (=O) include, but are not limited to, carbocyclics such as cyclopentanone and cyclohexanone; heterocyclics, such as pyrone, pyrrolidone, pyrazolone, pyrazolinone, piperidone, piperidinedione, piperazinedione, and imidazolidone; cyclic anhydrides, including but not limited to maleic anhydride and

- succinic anhydride; cyclic carbonates, such as propylene carbonate; imides, including but not limited to, succinimide and maleimide; lactones (cyclic esters, -O-C(=O)- in a ring), including, but not limited to, β -propiolactone, γ -butyrolactone, δ -valerolactone, and ϵ -caprolactone; and lactams (cyclic amides, -NH-C(=O)- in a ring), including, but not limited to, β -propiolactam, γ -butyrolactam, δ -valerolactam, and ϵ -caprolactam.

- 10 Imino (imine): =NR, wherein R is an imino substituent, for example, hydrogen, C_{1-7} alkyl group, a C_{3-20} heterocyclyl group, or a C_{5-20} aryl group, preferably hydrogen or a C_{1-7} alkyl group. Examples of imino groups include, but are not limited to, =NH, =NMe, =NEt, and =NPh.

Formyl (carbaldehyde, carboxaldehyde): -C(=O)H.

- 15 Acyl (keto): -C(=O)R, wherein R is an acyl substituent, for example, a C_{1-7} alkyl group (also referred to as C_{1-7} alkylacyl or C_{1-7} alkanoyl), a C_{3-20} heterocyclyl group (also referred to as C_{3-20} heterocyclylacyl), or a C_{5-20} aryl group (also referred to as C_{5-20} arylacyl), preferably a C_{1-7} alkyl group. Examples of acyl groups include, but are not limited to, -C(=O)CH₃ (acetyl), -C(=O)CH₂CH₃ (propionyl), -C(=O)C(CH₃)₃ (20 butyryl), and -C(=O)Ph (benzoyl, phenone).

Acylhalide (haloformyl, halocarbonyl): -C(=O)X, wherein X is -F, -Cl, -Br, or -I, preferably -Cl, -Br, or -I.

- 25 Carboxy (carboxylic acid): -COOH.

- Ester (carboxylate, carboxylic acid ester, oxycarbonyl): -C(=O)OR, wherein R is an ester substituent, for example, a C_{1-7} alkyl group, a C_{3-20} heterocyclyl group, or a C_{5-20} aryl group, preferably a C_{1-7} alkyl group. Examples of ester groups include, 30 but are not limited to, -C(=O)OCH₃, -C(=O)OCH₂CH₃, -C(=O)OC(CH₃)₃, and -C(=O)OPh.

Acyloxy (reverse ester): $-\text{OC}(=\text{O})\text{R}$, wherein R is an acyloxy substituent, for example, a C_{1-7} alkyl group, a C_{3-20} heterocyclyl group, or a C_{5-20} aryl group, preferably a C_{1-7} alkyl group. Examples of acyloxy groups include, but are not limited to, $-\text{OC}(=\text{O})\text{CH}_3$ (acetoxy), $-\text{OC}(=\text{O})\text{CH}_2\text{CH}_3$, $-\text{OC}(=\text{O})\text{C}(\text{CH}_3)_3$,

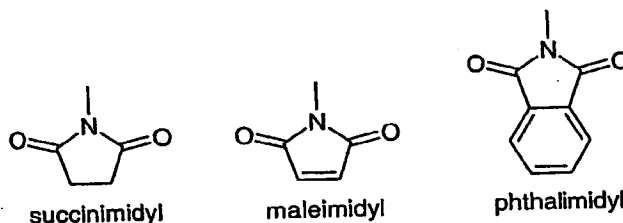
5 $-\text{OC}(=\text{O})\text{Ph}$, and $-\text{OC}(=\text{O})\text{CH}_2\text{Ph}$.

Amido (carbamoyl, carbamyl, aminocarbonyl, carboxamide): $-\text{C}(=\text{O})\text{NR}^1\text{R}^2$, wherein R^1 and R^2 are independently amino substituents, as defined for amino groups. Examples of amido groups include, but are not limited to, $-\text{C}(=\text{O})\text{NH}_2$,
10 $-\text{C}(=\text{O})\text{NHCH}_3$, $-\text{C}(=\text{O})\text{NH}(\text{CH}_3)_2$, $-\text{C}(=\text{O})\text{NHCH}_2\text{CH}_3$, and $-\text{C}(=\text{O})\text{N}(\text{CH}_2\text{CH}_3)_2$, as well as amido groups in which R^1 and R^2 , together with the nitrogen atom to which they are attached, form a heterocyclic structure as in, for example, piperidinocarbonyl, morpholinocarbonyl, thiomorpholinocarbonyl, and piperazinocarbonyl.

15

Acylamido (acylamino): $-\text{NR}^1\text{C}(=\text{O})\text{R}^2$, wherein R^1 is an amide substituent, for example, a C_{1-7} alkyl group, a C_{3-20} heterocyclyl group, or a C_{5-20} aryl group, preferably a C_{1-7} alkyl group, and R^2 is an acyl substituent, for example, a C_{1-7} alkyl group, a C_{3-20} heterocyclyl group, or a C_{5-20} aryl group, preferably a C_{1-7} alkyl group.

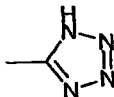
20 Examples of acylamido groups include, but are not limited to, $-\text{NHC}(=\text{O})\text{CH}_3$, $-\text{NHC}(=\text{O})\text{CH}_2\text{CH}_3$, and $-\text{NHC}(=\text{O})\text{Ph}$. R^1 and R^2 may together form a cyclic structure, as in, for example, succinimidyl, maleimidyl, and phthalimidyl:



25

Thioamido (thiocarbamyl): $-\text{C}(=\text{S})\text{NR}^1\text{R}^2$, wherein R^1 and R^2 are independently amino substituents, as defined for amino groups. Examples of amido groups include, but are not limited to, $-\text{C}(=\text{S})\text{NH}_2$, $-\text{C}(=\text{S})\text{NHCH}_3$, $-\text{C}(=\text{S})\text{NH}(\text{CH}_3)_2$, and $-\text{C}(=\text{S})\text{NHCH}_2\text{CH}_3$.

Tetrazolyl: a five membered aromatic ring having four nitrogen atoms and one carbon atom,



5

Amino: $-NR^1R^2$, wherein R^1 and R^2 are independently amino substituents, for example, hydrogen, a C_{1-7} alkyl group (also referred to as C_{1-7} alkylamino or di- C_{1-7} alkylamino), a C_{3-20} heterocyclyl group, or a C_{5-20} aryl group, preferably H or a C_{1-7} alkyl group, or, in the case of a "cyclic" amino group, R^1 and R^2 , taken

10 together with the nitrogen atom to which they are attached, form a heterocyclic ring having from 4 to 8 ring atoms. Examples of amino groups include, but are not limited to, $-NH_2$, $-NHCH_3$, $-NHCH(CH_3)_2$, $-N(CH_3)_2$, $-N(CH_2CH_3)_2$, and $-NHPh$. Examples of cyclic amino groups include, but are not limited to, aziridino, azetidino, piperidino, piperazino, morpholino, and thiomorpholino.

15

Nitro: $-NO_2$.

Nitroso: $-NO$.

20 Azido: $-N_3$.

Cyano (nitrile, carbonitrile): $-CN$.

Isocyano: $-NC$.

25

Cyanato: $-OCN$.

Isocyanato: $-NCO$.

30 Thiocyano (thiocyanato): $-SCN$.

Isothiocyano (isothiocyanato): $-NCS$.

Sulphydryl (thiol, mercapto): -SH.

- 5 Thioether (sulfide): -SR, wherein R is a thioether substituent, for example, a C₁₋₇alkyl group (also referred to as a C₁₋₇alkylthio group), a C₃₋₂₀heterocyclyl group, or a C₅₋₂₀aryl group, preferably a C₁₋₇alkyl group. Examples of C₁₋₇alkylthio groups include, but are not limited to, -SCH₃ and -SCH₂CH₃.

Sulfonic acid (sulfo): -S(=O)₂OH.

10

Sulfonate (sulfonic acid ester): -S(=O)₂OR, wherein R is a sulfonate substituent, for example, a C₁₋₇alkyl group, a C₃₋₂₀heterocyclyl group, or a C₅₋₂₀aryl group, preferably a C₁₋₇alkyl group. Examples of sulfonate groups include, but are not limited to, -S(=O)₂OCH₃ and -S(=O)₂OCH₂CH₃.

15

Sulfone (sulfonyl): -S(=O)₂R, wherein R is a sulfone substituent, for example, a C₁₋₇alkyl group, a C₃₋₂₀heterocyclyl group, or a C₅₋₂₀aryl group, preferably a C₁₋₇alkyl group. Examples of sulfone groups include, but are not limited to, -S(=O)₂CH₃ (methanesulfonyl, mesyl), -S(=O)₂CF₃, -S(=O)₂CH₂CH₃, and 4-

20 methylphenylsulfonyl (tosyl).

Sulfonyloxy: -OS(=O)₂R, wherein R is a sulfonyloxy substituent, for example, a C₁₋₇alkyl group, a C₃₋₂₀heterocyclyl group, or a C₅₋₂₀aryl group, preferably a C₁₋₇alkyl group. Examples of sulfonyloxy groups include, but are not limited to,

25 -OS(=O)₂CH₃ and -OS(=O)₂CH₂CH₃.

Sulfinyloxy: -OS(=O)R, wherein R is a sulfinyloxy substituent, for example, a C₁₋₇alkyl group, a C₃₋₂₀heterocyclyl group, or a C₅₋₂₀aryl group, preferably a C₁₋₇alkyl group. Examples of sulfinyloxy groups include, but are not limited to,

30 -OS(=O)CH₃ and -OS(=O)CH₂CH₃.

Sulfamino: $-\text{NR}^1\text{S}(=\text{O})_2\text{OH}$, wherein R^1 is an amino substituent, as defined for amino groups. Examples of sulfamino groups include, but are not limited to, $-\text{NHS}(=\text{O})_2\text{OH}$ and $-\text{N}(\text{CH}_3)\text{S}(=\text{O})_2\text{OH}$.

- 5 Sulfonamino: $-\text{NR}^1\text{S}(=\text{O})_2\text{R}$, wherein R^1 is an amino substituent, as defined for amino groups, and R is a sulfonamino substituent, for example, a C_{1-7} alkyl group, a C_{3-20} heterocyclyl group, or a C_{5-20} aryl group, preferably a C_{1-7} alkyl group. Examples of sulfonamino groups include, but are not limited to, $-\text{NHS}(=\text{O})_2\text{CH}_3$ and $-\text{N}(\text{CH}_3)\text{S}(=\text{O})_2\text{C}_6\text{H}_5$.

10

Sulfinamino: $-\text{NR}^1\text{S}(=\text{O})\text{R}$, wherein R^1 is an amino substituent, as defined for amino groups, and R is a sulfinamino substituent, for example, a C_{1-7} alkyl group, a C_{3-20} heterocyclyl group, or a C_{5-20} aryl group, preferably a C_{1-7} alkyl group. Examples of sulfinamino groups include, but are not limited to, $-\text{NHS}(=\text{O})\text{CH}_3$ and

- 15 $-\text{N}(\text{CH}_3)\text{S}(=\text{O})\text{C}_6\text{H}_5$.

Sulfamyl: $-\text{S}(=\text{O})\text{NR}^1\text{R}^2$, wherein R^1 and R^2 are independently amino substituents, as defined for amino groups. Examples of sulfamyl groups include, but are not limited to, $-\text{S}(=\text{O})\text{NH}_2$, $-\text{S}(=\text{O})\text{NH}(\text{CH}_3)$, $-\text{S}(=\text{O})\text{N}(\text{CH}_3)_2$, $-\text{S}(=\text{O})\text{NH}(\text{CH}_2\text{CH}_3)$,

- 20 $-\text{S}(=\text{O})\text{N}(\text{CH}_2\text{CH}_3)_2$, and $-\text{S}(=\text{O})\text{NHPh}$.

Sulfonamido: $-\text{S}(=\text{O})_2\text{NR}^1\text{R}^2$, wherein R^1 and R^2 are independently amino substituents, as defined for amino groups. Examples of sulfonamido groups include, but are not limited to, $-\text{S}(=\text{O})_2\text{NH}_2$, $-\text{S}(=\text{O})_2\text{NH}(\text{CH}_3)$, $-\text{S}(=\text{O})_2\text{N}(\text{CH}_3)_2$,

- 25 $-\text{S}(=\text{O})_2\text{NH}(\text{CH}_2\text{CH}_3)$, $-\text{S}(=\text{O})_2\text{N}(\text{CH}_2\text{CH}_3)_2$, and $-\text{S}(=\text{O})_2\text{NHPh}$.

- As mentioned above, a C_{1-7} alkyl group may be substituted with, for example, hydroxy (also referred to as a C_{1-7} hydroxyalkyl group), C_{1-7} alkoxy (also referred to as a C_{1-7} alkoxyalkyl group), amino (also referred to as a C_{1-7} aminoalkyl group),
- 30 halo (also referred to as a C_{1-7} haloalkyl group), carboxy (also referred to as a C_{1-7} carboxyalkyl group), and C_{5-20} aryl (also referred to as a C_{5-20} aryl- C_{1-7} alkyl group).

- Similarly, a C₅₋₂₀aryl group may be substituted with, for example, hydroxy (also referred to as a C₅₋₂₀hydroxyaryl group), halo (also referred to as a C₅₋₂₀haloaryl group), amino (also referred to as a C₅₋₂₀aminoaryl group, e.g., as in aniline), C₁₋₇alkyl (also referred to as a C₁₋₇alkyl-C₅₋₂₀aryl group, e.g., as in toluene), and
- 5 C₁₋₇alkoxy (also referred to as a C₁₋₇alkoxy-C₅₋₂₀aryl group, e.g., as in anisole).

These and other specific examples of such substituted groups are also discussed below.

- 10 C₁₋₇haloalkyl group: The term "C₁₋₇haloalkyl group," as used herein, pertains to a C₁₋₇alkyl group in which at least one hydrogen atom (e.g., 1, 2, 3) has been replaced with a halogen atom (e.g., F, Cl, Br, I). If more than one hydrogen atom has been replaced with a halogen atom, the halogen atoms may independently be the same or different. Every hydrogen atom may be replaced with a halogen
- 15 atom, in which case the group may conveniently be referred to as a C₁₋₇perhaloalkyl group." Examples of C₁₋₇haloalkyl groups include, but are not limited to, -CF₃, -CHF₂, -CH₂F, -CCl₃, -CBr₃, -CH₂CH₂F, -CH₂CHF₂, and -CH₂CF₃.
- 20 C₁₋₇hydroxyalkyl: The term "C₁₋₇hydroxyalkyl group," as used herein, pertains to a C₁₋₇alkyl group in which at least one hydrogen atom has been replaced with a hydroxy group. Examples of C₁₋₇hydroxyalkyl groups include, but are not limited to, -CH₂OH, -CH₂CH₂OH, and -CH(OH)CH₂OH.
- 25 C₁₋₇carboxyalkyl: The term "C₁₋₇carboxyalkyl group," as used herein, pertains to a C₁₋₇alkyl group in which at least one hydrogen atom has been replaced with a carboxy group. Examples of C₁₋₇carboxyalkyl groups include, but are not limited to, -CH₂COOH and -CH₂CH₂COOH.
- 30 C₁₋₇aminoalkyl: The term "C₁₋₇aminoalkyl group," as used herein, pertains to a C₁₋₇alkyl group in which at least one hydrogen atom has been replaced with an amino group. Examples of C₁₋₇aminoalkyl groups include, but are not limited to, -CH₂NH₂, -CH₂CH₂NH₂, and -CH₂CH₂N(CH₃)₂.

C₁₋₇alkyl-C₅₋₂₀aryl: The term "C₁₋₇alkyl-C₅₋₂₀aryl," as used herein, describes certain C₅₋₂₀aryl groups which have been substituted with a C₁₋₇alkyl group. Examples of such groups include, but are not limited to, tolyl (as in toluene), xylyl (as in xylene), mesityl (as in mesitylene), styryl (as in styrene), and cumenyl (as in cumene).

C₅₋₂₀aryl-C₁₋₇alkyl: The term "C₅₋₂₀aryl-C₁₋₇alkyl," as used herein, describes certain C₁₋₇alkyl groups which have been substituted with a C₅₋₂₀aryl group. Examples of such groups include, but are not limited to, benzyl (phenylmethyl), tolylmethyl, phenylethyl, and triphenylmethyl (trityl).

C₅₋₂₀haloaryl: The term "C₅₋₂₀haloaryl," as used herein, describes certain C₅₋₂₀aryl groups which have been substituted with one or more halo groups. Examples of such groups include, but are not limited to, halophenyl (e.g., fluorophenyl, chlorophenyl, bromophenyl, or iodophenyl, whether ortho-, meta-, or para-substituted), dihalophenyl, trihalophenyl, tetrahalophenyl, and pentahalophenyl.

Bidentate Substituents

Some substituents are bidentate, that is, have two points for covalent attachment. For example, a bidentate group may be covalently bound to two different atoms on two different groups, thereby acting as a linker therebetween. Alternatively, a bidentate group may be covalently bound to two different atoms on the same group, thereby forming, together with the two atoms to which it is attached (and any intervening atoms, if present) a cyclic or ring structure. In this way, the bidentate substituent may give rise to a heterocyclic group/compound and/or an aromatic group/compound. Typically, the ring has from 3 to 8 ring atoms, which ring atoms are carbon or divalent heteroatoms (e.g., boron, silicon, nitrogen, phosphorus, oxygen, and sulfur, typically nitrogen, oxygen, and sulfur), and wherein the bonds between said ring atoms are single or double bonds, as permitted by the valencies of the ring atoms. Typically, the bidentate group is covalently bound to vicinal atoms, that is, adjacent atoms, in the parent group.

C₁₋₇alkylene: The term "C₁₋₇alkylene," as used herein, pertains to a bidentate moiety obtained by removing two hydrogen atoms, either both from the same carbon atom, or one from each of two different carbon atoms, of a C₁₋₇hydrocarbon compound having from 1 to 7 carbon atoms, which may be aliphatic or alicyclic, or a combination thereof, and which may be saturated, partially unsaturated, or fully unsaturated.

Examples of linear saturated C₁₋₇alkylene groups include, but are not limited to, - (CH₂)_n- where n is an integer from 1 to 7, for example, -CH₂- (methylene), - CH₂CH₂- (ethylene), -CH₂CH₂CH₂- (propylene), and -CH₂CH₂CH₂CH₂- (butylene).

Examples of branched saturated C₁₋₇alkylene groups include, but are not limited to, -CH(CH₃)-, -CH(CH₃)CH₂-, -CH(CH₃)CH₂CH₂-, -CH(CH₃)CH₂CH₂CH₂-, -CH₂CH(CH₃)CH₂-, -CH₂CH(CH₃)CH₂CH₂-, -CH(CH₂CH₃)-, -CH(CH₂CH₃)CH₂-, and -CH₂CH(CH₂CH₃)CH₂-.

Examples of linear partially unsaturated C₁₋₇alkylene groups include, but are not limited to, -CH=CH- (vinylene), -CH=CH-CH₂-, -CH=CH-CH₂-CH₂-, - CH=CH-CH₂-CH₂-CH₂-, -CH=CH-CH=CH-, -CH=CH-CH=CH-CH₂-, -CH=CH- CH=CH-CH₂-CH₂-, -CH=CH-CH₂-CH=CH-, and -CH=CH-CH₂-CH₂-CH=CH-.

Examples of branched partially unsaturated C₁₋₇alkylene groups include, but are not limited to, -C(CH₃)=CH-, -C(CH₃)=CH-CH₂-, and -CH=CH-CH(CH₃)-.

Examples of alicyclic saturated C₁₋₇alkylene groups include, but are not limited to, cyclopentylene (e.g., cyclopent-1,3-ylene), and cyclohexylene (e.g., cyclohex-1,4-ylene).

Examples of alicyclic partially unsaturated C₁₋₇alkylene groups include, but are not limited to, cyclopentenylene (e.g., 4-cyclopenten-1,3-ylene), cyclohexenylene (e.g., 2-cyclohexen-1,4-ylene, 3-cyclohexen-1,2-ylene, 2,5-cyclohexadien-1,4-ylene).

C₅₋₂₀arylene: The term "C₅₋₂₀arylene," as used herein, pertains to a bidentate moiety obtained by removing two hydrogen atoms, one from each of two different ring atoms of a C₅₋₂₀aromatic compound, said compound having one ring, or two or more rings (e.g., fused), and having from 5 to 20 ring atoms, and wherein at least one of said ring(s) is an aromatic ring. Preferably, each ring has from 5 to 7 ring atoms.

The ring atoms may be all carbon atoms, as in "carboarylene groups," in which case the group may conveniently be referred to as a "C₅₋₂₀carboarylene" group.

10

Alternatively, the ring atoms may include one or more heteroatoms, including but not limited to oxygen, nitrogen, and sulfur, as in "heteroarylene groups." In this case, the group may conveniently be referred to as a "C₅₋₂₀heteroarylene" group, wherein "C₅₋₂₀" denotes ring atoms, whether carbon atoms or heteroatoms.

15 Preferably, each ring has from 5 to 7 ring atoms, of which from 0 to 4 are ring heteroatoms.

Examples of C₅₋₂₀arylene groups which do not have ring heteroatoms (i.e., C₅₋₂₀carboarylene groups) include, but are not limited to, those derived from benzene (i.e., phenyl) (C₆), naphthalene (C₁₀), anthracene (C₁₄), phenanthrene (C₁₄), and pyrene (C₁₆).

Examples of C₅₋₂₀heteroarylene groups include, but are not limited to, C₅heteroarylene groups derived from furan (oxole), thiophene (thiole), pyrrole (azole), imidazole (1,3-diazole), pyrazole (1,2-diazole), triazole, oxazole, isoxazole, thiazole, isothiazole, oxadiazole, and oxatriazole; and C₆heteroarylene groups derived from isoxazine, pyridine (azine), pyridazine (1,2-diazine), pyrimidine (1,3-diazine; e.g., cytosine, thymine, uracil), pyrazine (1,4-diazine), triazine, tetrazole, and oxadiazole (furazan).

30 C₅₋₂₀Arylene-C₁₋₇alkylene: The term "C₅₋₂₀arylene-C₁₋₇alkylene," as used herein, pertains to a bidentate moiety comprising a C₅₋₂₀arylene moiety, -Arylene-, linked to a C₁₋₇alkylene moiety, -Alkylene-, that is, -Arylene-Alkylene-.

Examples of C₅₋₂₀arylene-C₁₋₇alkylene groups include, but are not limited to, phenylene-methylene, phenylene-ethylene, phenylene-propylene, and phenylene-ethenylene (also known as phenylene-vinylene).

- 5 C₅₋₂₀Alkylene-C₁₋₇arylene: The term "C₅₋₂₀alkylene-C₁₋₇arylene," as used herein, pertains to a bidentate moiety comprising a C₅₋₂₀alkylene moiety, -Alkylene-, linked to a C₁₋₇arylene moiety, -Arylene-, that is, -Alkylene-Arylene-.

- 10 Examples of C₅₋₂₀alkylene-C₁₋₇arylene groups include, but are not limited to, methylene-phenylene, ethylene-phenylene, propylene-phenylene, and ethenylene-phenylene (also known as vinylene-phenylene).

- 15 Included in the above are the well known ionic, salt, solvate (e.g., hydrate), and protected forms of these substituents. For example, a reference to carboxylic acid (-COOH) also includes carboxylate (-COO⁻). Similarly, a reference to an amino group includes a salt, for example, a hydrochloride salt, of the amino group. A reference to a hydroxyl group also includes conventional protected forms of a hydroxyl group. Similarly, a reference to an amino group also includes conventional protected forms of an amino group.

20

Acronyms

- 25 For convenience, many chemical moieties are represented herein using well known abbreviations, including but not limited to, methyl (Me), ethyl (Et), n-propyl (nPr), iso-propyl (iPr), n-butyl (nBu), tert-butyl (tBu), n-hexyl (nHex), cyclohexyl (cHex), phenyl (Ph), biphenyl (biPh), benzyl (Bn), naphthyl (naph), methoxy (MeO), ethoxy (EtO), benzoyl (Bz), and acetyl (Ac).

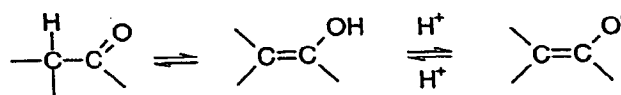
- 30 For convenience, many chemical compounds are represented herein using well known abbreviations, including but not limited to, methanol (MeOH), ethanol (EtOH), iso-propanol (i-PrOH), methyl ethyl ketone (MEK), acetic acid (AcOH), dichloromethane (methylene chloride, DCM), trifluoroacetic acid (TFA), dimethylformamide (DMF), and tetrahydrofuran (THF).

Isomers, Salts, Hydrates, Protected Forms, and Prodrugs

5 A certain compound may exist in one or more particular geometric, optical, enantiomeric, diastereomeric, epimeric, stereoisomeric, tautomeric, conformational, or anomeric forms, including but not limited to, cis- and trans-forms; E- and Z-forms; c-, t-, and r- forms; endo- and exo-forms; R-, S-, and meso-forms; D- and L-forms; (+) and (-) forms; keto-, enol-, and enolate-forms; syn- and anti-forms; synclinal- and anticlinal-forms; α - and β -forms; axial and
10 equatorial forms; boat-, chair-, twist-, envelope-, and halfchair-forms; and combinations thereof, hereinafter collectively referred to as "isomers" (or "isomeric forms").

Note that, except as discussed below for tautomeric forms, specifically excluded
15 from the term "isomers," as used herein, are structural (or constitutional) isomers (i.e., isomers which differ in the connections between atoms rather than merely by the position of atoms in space). For example, a reference to a methoxy group, -OCH₃, is not to be construed as a reference to its structural isomer, a hydroxymethyl group, -CH₂OH. Similarly, a reference to ortho-chlorophenyl is not
20 to be construed as a reference to its structural isomer, meta-chlorophenyl. However, a reference to a class of structures may well include structurally isomeric forms falling within that class (e.g., C₁₋₇alkyl includes n-propyl and isopropyl; butyl includes n-, iso-, sec-, and tert-butyl; methoxyphenyl includes ortho-, meta-, and para-methoxyphenyl).

25 The above exclusion does not pertain to tautomeric forms, for example, keto-, enol-, and enolate-forms, as in, for example, the following tautomeric pairs: keto/enol (illustrated below), imine/enamine, amide/imino alcohol, amidine/amidine, nitroso/oxime, thioketone/enethiol, N-nitroso/hydroxyazo, and
30 nitro/aci-nitro.



Note that specifically included in the term "isomer" are compounds with one or more isotopic substitutions. For example, H may be in any isotopic form, including ^1H , ^2H (D), and ^3H (T); C may be in any isotopic form, including ^{12}C , ^{13}C , and ^{14}C ; O may be in any isotopic form, including ^{16}O and ^{18}O ; and the like.

5

Unless otherwise specified, a reference to a particular compound includes all such isomeric forms, including racemic and other mixtures thereof. Methods for the preparation (e.g., asymmetric synthesis) and separation (e.g., fractional crystallisation and chromatographic means) of such isomeric forms are either
10 known in the art or are readily obtained by adapting the methods taught herein in a known manner.

15

Unless otherwise specified, a reference to a particular compound also includes ionic, salt, solvate (e.g., hydrate), protected forms, and prodrugs thereof, for example, as discussed below.

20

It may be convenient or desirable to prepare, purify, and/or handle a corresponding salt of the active compound, for example, a pharmaceutically-acceptable salt. Examples of pharmaceutically acceptable salts are discussed in Berge et al., 1977, "Pharmaceutically Acceptable Salts," J. Pharm. Sci., Vol. 66, pp. 1-19.

25

For example, if the compound is anionic, or has a functional group which may be anionic (e.g., $-\text{COOH}$ may be $-\text{COO}^-$), then a salt may be formed with a suitable cation. Examples of suitable inorganic cations include, but are not limited to, alkali metal ions such as Na^+ and K^+ , alkaline earth cations such as Ca^{2+} and Mg^{2+} , and other cations such as Al^{3+} . Examples of suitable organic cations include, but are not limited to, ammonium ion (i.e., NH_4^+) and substituted ammonium ions (e.g., NH_3R^+ , NH_2R_2^+ , NHR_3^+ , NR_4^+). Examples of some suitable substituted
30 ammonium ions are those derived from: ethylamine, diethylamine, ethylenediamine, ethanolamine, diethanolamine, piperazine. An example of a common quaternary ammonium ion is $\text{N}(\text{CH}_3)_4^+$.

If the compound is cationic, or has a functional group which may be cationic (e.g., $-\text{NH}_2$ may be $-\text{NH}_3^+$), then a salt may be formed with a suitable anion. Examples of suitable inorganic anions include, but are not limited to, those derived from the following inorganic acids: hydrochloric, hydrobromic, hydroiodic, sulfuric, sulfurous, nitric, nitrous, phosphoric, and phosphorous. Examples of suitable organic anions include, but are not limited to, anions from the following organic acids: acetic, propionic, succinic, glycolic, stearic, lactic, malic, tartaric, citric, ascorbic, maleic, hydroxymaleic, phenylacetic, glutamic, benzoic, salicylic, sulfanilic, 2-acetoxybenzoic, fumaric, toluenesulfonic, methanesulfonic, ethanesulfonic, ethane disulfonic, oxalic, isethionic, and valeric.

It may be convenient or desirable to prepare, purify, and/or handle a corresponding solvate of the active compound. The term "solvate" is used herein in the conventional sense to refer to a complex of solute (e.g., active compound, salt of active compound) and solvent. If the solvent is water, the solvate may be conveniently referred to as a hydrate, for example, a mono-hydrate, a di-hydrate, a tri-hydrate, etc.

It may be convenient or desirable to prepare, purify, and/or handle the active compound in a chemically protected form. The term "chemically protected form," as used herein, pertains to a compound in which one or more reactive functional groups are protected from undesirable chemical reactions, that is, are in the form of a protected or protecting group (also known as a masked or masking group). By protecting a reactive functional group, reactions involving other unprotected reactive functional groups can be performed, without affecting the protected group; the protecting group may be removed, usually in a subsequent step, without substantially affecting the remainder of the molecule. See, for example, Protective Groups in Organic Synthesis (T. Green and P. Wuts, Wiley, 1991), and Protective Groups in Organic Synthesis (T. Green and P. Wuts; 3rd Edition; John Wiley and Sons, 1999).

For example, a hydroxy group may be protected as an ether ($-\text{OR}$) or an ester ($-\text{OC}(=\text{O})\text{R}$), for example, as: a t-butyl ether; a benzyl, benzhydryl

(diphenylmethyl), or trityl (triphenylmethyl) ether; a trimethylsilyl or t-butyl dimethylsilyl ether; or an acetyl ester ($-\text{OC}(=\text{O})\text{CH}_3$, $-\text{OAc}$).

For example, an aldehyde or ketone group may be protected as an acetal or ketal, respectively, in which the carbonyl group ($>\text{C}=\text{O}$) is converted to a diether ($>\text{C}(\text{OR})_2$), by reaction with, for example, a primary alcohol. The aldehyde or ketone group is readily regenerated by hydrolysis using a large excess of water in the presence of acid.

For example, an amine group may be protected, for example, as an amide ($-\text{NRCO}-\text{R}$) or a urethane ($-\text{NRCO}-\text{OR}$), for example, as: a methyl amide ($-\text{NHCO}-\text{CH}_3$); a benzyloxy amide ($-\text{NHCO}-\text{OCH}_2\text{C}_6\text{H}_5$, $-\text{NH}-\text{Cbz}$); as a t-butoxy amide ($-\text{NHCO}-\text{OC}(\text{CH}_3)_3$, $-\text{NH}-\text{Boc}$); a 2-biphenyl-2-propoxy amide ($-\text{NHCO}-\text{OC}(\text{CH}_3)_2\text{C}_6\text{H}_4\text{C}_6\text{H}_5$, $-\text{NH}-\text{Bpoc}$), as a 9-fluorenylmethoxy amide ($-\text{NH}-\text{Fmoc}$), as a 6-nitroveratryloxy amide ($-\text{NH}-\text{Nvoc}$), as a 2-trimethylsilylethyloxy amide ($-\text{NH}-\text{Teoc}$), as a 2,2,2-trichloroethyloxy amide ($-\text{NH}-\text{Troc}$), as an allyloxy amide ($-\text{NH}-\text{Alloc}$), as a 2-(phenylsulfonyl)ethyloxy amide ($-\text{NH}-\text{Psec}$); or, in suitable cases (e.g., cyclic amines), as a nitroxide radical ($>\text{N}-\text{O}\cdot$).

20

For example, a carboxylic acid group may be protected as an ester or an amide, for example, as: a benzyl ester; a t-butyl ester; a methyl ester; or a methyl amide.

For example, a thiol group may be protected as a thioether ($-\text{SR}$), for example, as: a benzyl thioether; an acetamidomethyl ether ($-\text{S}-\text{CH}_2\text{NHC}(=\text{O})\text{CH}_3$).

It may be convenient or desirable to prepare, purify, and/or handle the active compound in the form of a prodrug. The term "prodrug," as used herein, pertains to a compound which, when metabolised, yields the desired active compound. Typically, the prodrug is inactive, or less active than the active compound, but may provide advantageous handling, administration, or metabolic properties. For example, some prodrugs are esters of the active compound; during metabolism, the ester group is cleaved to yield the active drug. Also, some prodrugs are

activated enzymatically to yield the active compound, or a compound which; upon further chemical reaction, yields the active compound. For example, the prodrug may be a sugar derivative or other glycoside conjugate, or may be an amino acid ester derivative.

5

Synthesis

Several methods for the chemical synthesis of compounds of the present invention are described herein. These methods may be modified and/or adapted in known ways in order to facilitate the synthesis of additional compounds within the scope of the present invention.

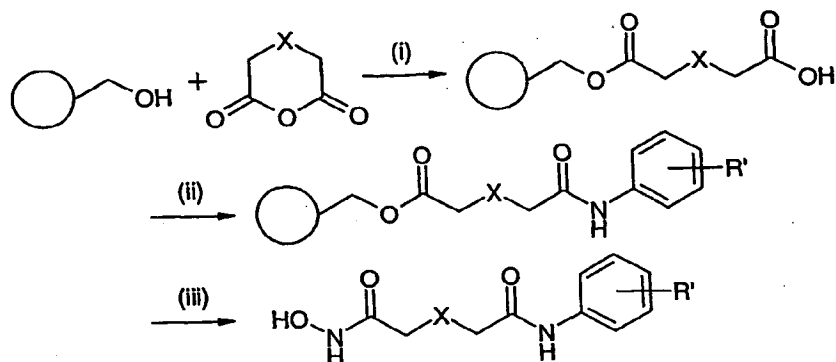
The compounds of the present invention may be prepared, for example, by the methods described herein, or by adapting these or other well known methods in well known ways.

In one method, a suitable resin with pendant hydroxy groups is reacted with a suitable carboxylic acid or anhydride, to form a pendant group which is linked to the solid support via an ester group, and which has a terminal carboxylic acid group. The terminal carboxylic acid group is then reacted with a suitable aryl amine (e.g., NH_2R), for example, an aniline, to give a terminal aryl amide. The resulting compound is then cleaved from the resin using hydroxylamine to give the desired carbamic acid.

One example of this approach is illustrated below, wherein the resin is ArgoGelJ-OH resin, X is -O- or -S-, and the reaction conditions are as follows: (i) DCM/pyridine (1:1, v/v), DMAP, room temperature, 16 hours; (ii) NH_2R , HOBT, TBTU, DIPEA, NMP, room temperature, 16 hours; and (iii) 50% NH_2OH , dioxane, room temperature, 48 hours.

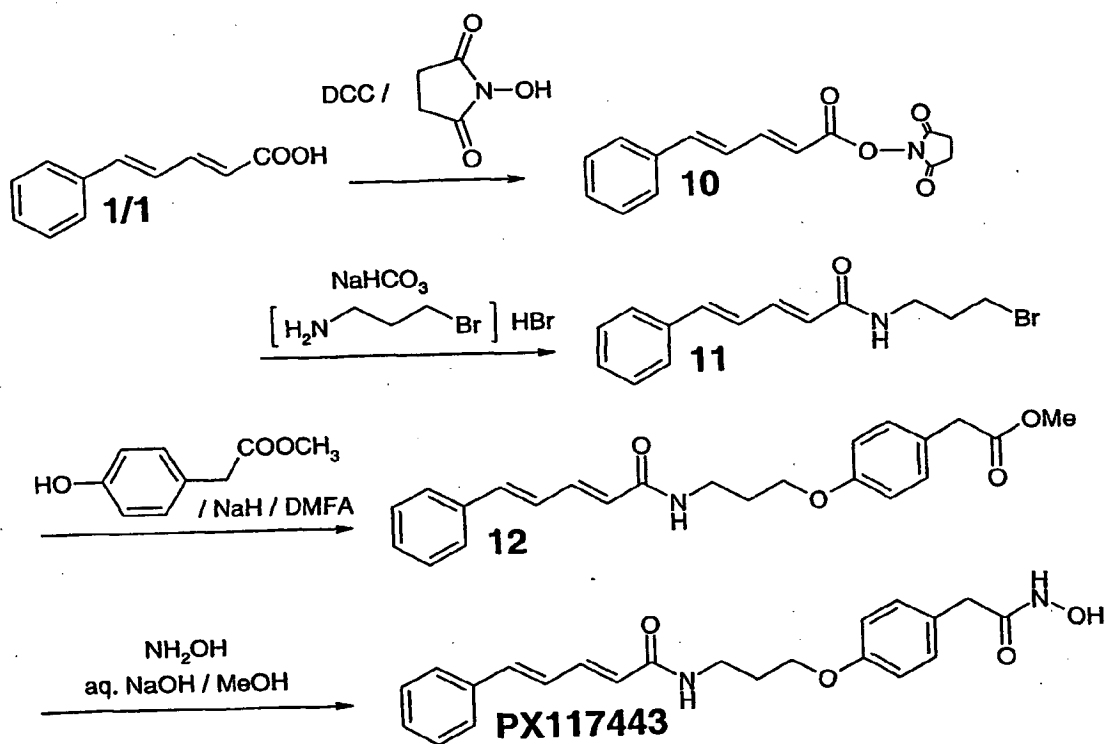
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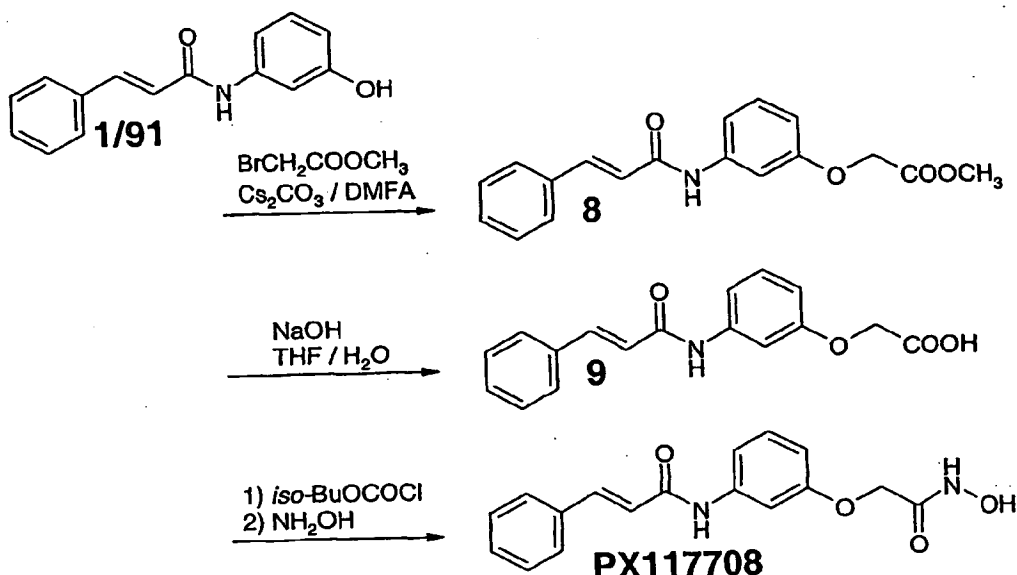
Scheme 1



Additional methods for the synthesis of compounds of the present invention are illustrated below and are exemplified in the examples below.

5

Scheme 2

Scheme 3Uses

5

The present invention provides active compounds which are capable of inhibiting HDAC (for example, inhibiting HDAC activity, inhibiting formation of HDAC complexes, inhibiting activity of HDAC complexes), as well as methods of inhibiting HDAC activity, comprising contacting a cell with an effective amount of an active compound, whether *in vitro* or *in vivo*.

10

The term "active," as used herein, pertains to compounds which are capable of inhibiting HDAC activity, and specifically includes both compounds with intrinsic activity (drugs) as well as prodrugs of such compounds, which prodrugs may themselves exhibit little or no intrinsic activity.

15

One of ordinary skill in the art is readily able to determine whether or not a candidate compound is active, that is, capable of inhibiting HDAC activity. For example, assays which may conveniently be used to assess the inhibition offered by a particular compound are described in the examples below.

20

- For example, a sample of cells (e.g., from a tumour) may be grown *in vitro* and a candidate compound brought into contact with the cells, and the effect of the compound on those cells observed. As examples of "effect," the morphological status of the cells may be determined (e.g., alive or dead), or the expression levels of genes regulated by HDAC. Where the candidate compound is found to exert an influence on the cells, this may be used as a prognostic or diagnostic marker of the efficacy of the compound in methods of treating a patient carrying cells of the same type (e.g., the tumour or a tumour of the same cellular type).
- 10 In one aspect, the present invention provides antiproliferative agents. The term "antiproliferative agent" as used herein, pertains to a compound which treats a proliferative condition (i.e., a compound which is useful in the treatment of a proliferative condition).
- 15 The terms "cell proliferation," "proliferative condition," "proliferative disorder," and "proliferative disease," are used interchangeably herein and pertain to an unwanted or uncontrolled cellular proliferation of excessive or abnormal cells which is undesired, such as, neoplastic or hyperplastic growth, whether *in vitro* or *in vivo*. Examples of proliferative conditions include, but are not limited to,
- 20 pre-malignant and malignant cellular proliferation, including but not limited to, malignant neoplasms and tumours, cancers, leukemias, psoriasis, bone diseases, fibroproliferative disorders (e.g., of connective tissues), and atherosclerosis. Any type of cell may be treated, including but not limited to, lung, colon, breast, ovarian, prostate, liver, pancreas, brain, and skin.
- 25 Antiproliferative compounds of the present invention have application in the treatment of cancer, and so the present invention further provides anticancer agents. The term "anticancer agent" as used herein, pertains to a compound which treats a cancer (i.e., a compound which is useful in the treatment of a
- 30 cancer). The anti-cancer effect may arise through one or more mechanisms, including but not limited to, the regulation of cell proliferation, the inhibition of angiogenesis (the formation of new blood vessels), the inhibition of metastasis (the spread of a tumour from its origin), the inhibition of invasion (the spread of

tumour cells into neighbouring normal structures), or the promotion of apoptosis (programmed cell death).

5 The compounds of the present invention may also be used in the treatment of conditions which are known to be mediated by HDAC, or which are known to be treated by HDAC inhibitors (such as, e.g., trichostatin A). Examples of such conditions include, but are not limited to, the following:

10 Cancer (see, e.g., Vigushin et al., 2001).

Psoriasis (see, e.g., Iavarone et al., 1999).

Fibroproliferative disorders (e.g., liver fibrosis) (see, e.g., Niki et al., 1999; Corneil et al., 1998).

15 Smooth muscle proliferative disorder (e.g., atherosclerosis, restenosis) (see, e.g., Kimura et al., 1994).

20 Neurodegenerative diseases (e.g., Alzheimer's, Parkinson's, Huntington's chorea, amyotrophic lateral sclerosis, spino-cerebellar degeneration) (see, e.g., Kuusisto et al., 2001).

Inflammatory disease (e.g., osteoarthritis, rheumatoid arthritis) (see, e.g., Dangond et al., 1998; Takahashi et al., 1996).

25 Diseases involving angiogenesis (e.g., cancer, rheumatoid arthritis, psoriasis, diabetic retinopathy) (see, e.g., Kim et al., 2001).

30 Haematopoietic disorders (e.g., anaemia, sickle cell anaemia, thalassaemia) (see, e.g., McCaffrey et al., 1997).

Fungal infection (see, e.g., Bernstein et al., 2000; Tsuji et al., 1976).

Parasitic infection (e.g., malaria, trypanosomiasis, helminthiasis, protozoal infections (see, e.g., Andrews et al., 2000)).

Bacterial infection (see, e.g., Onishi et al., 1996).

5

Viral infection (see, e.g., Chang et al., 2000).

Conditions treatable by immune modulation (e.g., multiple sclerosis, autoimmune diabetes, lupus, atopic dermatitis, allergies, asthma, allergic rhinitis, inflammatory
10 bowel disease; and for improving grafting of transplants) (see, e.g., Dangond et al., 1998; Takahashi et al., 1996).

The invention further provides active compounds for use in a method of treatment of the human or animal body. Such a method may comprise administering to
15 such a subject a therapeutically-effective amount of an active compound, preferably in the form of a pharmaceutical composition.

The term "treatment," as used herein in the context of treating a condition, pertains generally to treatment and therapy, whether of a human or an animal
20 (e.g., in veterinary applications), in which some desired therapeutic effect is achieved, for example, the inhibition of the progress of the condition, and includes a reduction in the rate of progress, a halt in the rate of progress, amelioration of the condition, and cure of the condition. Treatment as a prophylactic measure is also included.

25

The term "therapeutically-effective amount," as used herein, pertains to that amount of an active compound, or a material, composition or dosage from comprising an active compound, which is effective for producing some desired therapeutic effect, commensurate with a reasonable benefit/risk ratio.

30

The term "treatment" includes combination treatments and therapies, in which two or more treatments or therapies are combined, for example, sequentially or simultaneously. Examples of treatments and therapies include, but are not limited

to, chemotherapy (the administration of active agents, including, e.g., drugs, antibodies (e.g., as in immunotherapy), prodrugs (e.g., as in photodynamic therapy, GDEPT, ADEPT, etc.); surgery; radiation therapy; and gene therapy.

- 5 The invention further provides the use of an active compound for the manufacture of a medicament, for example, for the treatment of a proliferative condition, as discussed above.

- 10 The invention further provides the use of an active compound for the manufacture of a medicament, for example, for the treatment of conditions which are known to be mediated by HDAC, or which are known to be treated by HDAC inhibitors (such as, e.g., trichostatin A), as discussed above.

- 15 The invention further provides a method for inhibiting HDAC in a cell comprising said cell with an effective amount of an active compound.

- 20 The invention further provides a method of treatment of the human or animal body, the method comprising administering to a subject in need of treatment a therapeutically-effective amount of an active compound, preferably in the form of a pharmaceutical composition.

Active compounds may also be used, as described above, in combination therapies, that is, in conjunction with other agents, for example, cytotoxic agents.

- 25 Active compounds may also be used as part of an *in vitro* assay, for example, in order to determine whether a candidate host is likely to benefit from treatment with the compound in question.

- 30 Active compounds may also be used as a standard, for example, in an assay, in order to identify other active compounds, other antiproliferative agents, etc.

The compounds of the present invention may also be used in methods of improving protein production by cultured cells (see, e.g., Furukawa et al., 1998).

Routes of Administration

5 The active compound or pharmaceutical composition comprising the active compound may be administered to a subject by any convenient route of administration, whether systemically/ peripherally or topically (i.e., at the site of desired action).

10 Routes of administration include, but are not limited to, oral (e.g., by ingestion); buccal; sublingual; transdermal (including, e.g., by a patch, plaster, etc.); transmucosal (including, e.g., by a patch, plaster, etc.); intranasal (e.g., by nasal spray); ocular (e.g., by eyedrops); pulmonary (e.g., by inhalation or insufflation therapy using, e.g., via an aerosol, e.g., through the mouth or nose); rectal (e.g., by suppository or enema); vaginal (e.g., by pessary); parenteral, for example, by
15 injection, including subcutaneous, intradermal, intramuscular, intravenous, intraarterial, intracardiac, intrathecal, intraspinal, intracapsular, subcapsular, intraorbital, intraperitoneal, intratracheal, subcuticular, intraarticular, subarachnoid, and intrasternal; by implant of a depot or reservoir, for example, subcutaneously or intramuscularly.

20

The Subject

The subject may be a prokaryote (e.g., bacteria) or a eukaryote (e.g., protoctista, fungi, plants, animals).

25

The subject may be a protoctista, an alga, or a protozoan.

The subject may be a plant, an angiosperm, a dicotyledon, a monocotyledon, a gymnosperm, a conifer, a ginkgo, a cycad, a fern, a horsetail, a clubmoss, a
30 liverwort, or a moss.

The subject may be an animal.

The subject may be a chordate, an invertebrate, an echinoderm (e.g., starfish, sea urchins, brittlestars), an arthropod, an annelid (segmented worms) (e.g., earthworms, lugworms, leeches), a mollusk (cephalopods (e.g., squids, octopi), pelecypods (e.g., oysters, mussels, clams), gastropods (e.g., snails, slugs)), a
5 nematode (round worms), a platyhelminthes (flatworms) (e.g., planarians, flukes, tapeworms), a cnidaria (e.g., jelly fish, sea anemones, corals), or a porifera (e.g., sponges).

The subject may be an arthropod, an insect (e.g., beetles, butterflies, moths), a
10 chilopoda (centipedes), a diplopoda (millipedes), a crustacean (e.g., shrimps, crabs, lobsters), or an arachnid (e.g., spiders, scorpions, mites).

The subject may be a chordate, a vertebrate, a mammal, a bird, a reptile (e.g., snakes, lizards, crocodiles), an amphibian (e.g., frogs, toads), a bony fish (e.g.,
15 salmon, plaice, eel, lungfish), a cartilaginous fish (e.g., sharks, rays), or a jawless fish (e.g., lampreys, hagfish).

The subject may be a mammal, a placental mammal, a marsupial (e.g., kangaroo, wombat), a monotreme (e.g., duckbilled platypus), a rodent (e.g., a guinea pig, a
20 hamster, a rat, a mouse), murine (e.g., a mouse), a lagomorph (e.g., a rabbit), avian (e.g., a bird), canine (e.g., a dog), feline (e.g., a cat), equine (e.g., a horse), porcine (e.g., a pig), ovine (e.g., a sheep), bovine (e.g., a cow), a primate, simian (e.g., a monkey or ape), a monkey (e.g., marmoset, baboon), an ape (e.g., gorilla, chimpanzee, orangutang, gibbon), or a human.

25 Furthermore, the subject may be any of its forms of development, for example, a spore, a seed, an egg, a larva, a pupa, or a foetus.

In one preferred embodiment, the subject is a human.

30

Formulations

While it is possible for the active ingredient to be administered alone, it is preferable to present it as a pharmaceutical composition (e.g., formulation)

5 comprising at least one active ingredient, as defined above, together with one or more pharmaceutically acceptable carriers, excipients, buffers, adjuvants, stabilisers, or other materials well known to those skilled in the art and optionally other therapeutic agents.

10 Thus, the present invention further provides pharmaceutical compositions, as defined above, and methods of making a pharmaceutical composition comprising admixing at least one active ingredient, as defined above, together with one or more pharmaceutically acceptable carriers, excipients, buffers, adjuvants, stabilisers, or other materials, as described herein.

15 The term "pharmaceutically acceptable" as used herein pertains to compounds, materials, compositions, and/or dosage forms which are, within the scope of sound medical judgement, suitable for use in contact with the tissues of a subject (e.g., human) without excessive toxicity, irritation, allergic response, or other
20 problem or complication, commensurate with a reasonable benefit/risk ratio. Each carrier, excipient, etc. must also be "acceptable" in the sense of being compatible with the other ingredients of the formulation.

The formulations may conveniently be presented in unit dosage form and may be
25 prepared by any methods well known in the art of pharmacy. Such methods include the step of bringing into association the active ingredient with the carrier which constitutes one or more accessory ingredients. In general, the formulations are prepared by uniformly and intimately bringing into association the active ingredient with liquid carriers or finely divided solid carriers or both, and then if
30 necessary shaping the product.

Formulations may be in the form of liquids, solutions, suspensions, emulsions, tablets, lozenges, granules, powders, capsules, cachets, pills, ampoules,

suppositories, pessaries, ointments, gels, pastes, creams, sprays, foams, lotions, oils, boluses, electuaries, or aerosols.

5 Formulations suitable for oral administration (e.g., by ingestion) may be presented as discrete units such as capsules, cachets or tablets, each containing a predetermined amount of the active ingredient; as a powder or granules; as a solution or suspension in an aqueous or non-aqueous liquid; or as an oil-in-water liquid emulsion or a water-in-oil liquid emulsion; as a bolus; as an electuary; or as a paste.

10 A tablet may be made by compression or moulding, optionally with one or more accessory ingredients. Compressed tablets may be prepared by compressing in a suitable machine the active ingredient in a free-flowing form such as a powder or granules, optionally mixed with a binder (e.g., povidone, gelatin, hydroxypropylmethyl cellulose), lubricant, inert diluent, preservative, disintegrant
15 (e.g., sodium starch glycolate, cross-linked povidone, cross-linked sodium carboxymethyl cellulose), surface-active or dispersing agent. Moulded tablets may be made by moulding in a suitable machine a mixture of the powdered compound moistened with an inert liquid diluent. The tablets may optionally be
20 coated or scored and may be formulated so as to provide slow or controlled release of the active ingredient therein using, for example, hydroxypropylmethyl cellulose in varying proportions to provide the desired release profile. Tablets may optionally be provided with an enteric coating, to provide release in parts of the gut other than the stomach.

25 Formulations suitable for topical administration (e.g., transdermal, intranasal, ocular, buccal, and sublingual) may be formulated as an ointment, cream, suspension, lotion, powder, solution, paste, gel, spray, aerosol, or oil. Alternatively, a formulation may comprise a patch or a dressing such as a
30 bandage or adhesive plaster impregnated with active ingredients and optionally one or more excipients or diluents.

Formulations suitable for topical administration in the mouth include lozenges comprising the active ingredient in a flavored basis, usually sucrose and acacia or tragacanth; pastilles comprising the active ingredient in an inert basis such as gelatin and glycerin, or sucrose and acacia; and mouthwashes comprising the active ingredient in a suitable liquid carrier.

Formulations suitable for topical administration to the eye also include eye drops wherein the active ingredient is dissolved or suspended in a suitable carrier, especially an aqueous solvent for the active ingredient.

Formulations suitable for nasal administration, wherein the carrier is a solid, include a coarse powder having a particle size, for example, in the range of about 20 to about 500 microns which is administered in the manner in which snuff is taken, i.e., by rapid inhalation through the nasal passage from a container of the powder held close up to the nose. Suitable formulations wherein the carrier is a liquid for administration as, for example, nasal spray, nasal drops, or by aerosol administration by nebuliser, include aqueous or oily solutions of the active ingredient.

Formulations suitable for topical administration via the skin include ointments, creams, and emulsions. When formulated in an ointment, the active ingredient may optionally be employed with either a paraffinic or a water-miscible ointment base. Alternatively, the active ingredients may be formulated in a cream with an oil-in-water cream base. If desired, the aqueous phase of the cream base may include, for example, at least about 30% w/w of a polyhydric alcohol, i.e., an alcohol having two or more hydroxyl groups such as propylene glycol, butane-1,3-diol, mannitol, sorbitol, glycerol and polyethylene glycol and mixtures thereof. The topical formulations may desirably include a compound which enhances absorption or penetration of the active ingredient through the skin or other affected areas. Examples of such dermal penetration enhancers include dimethylsulfoxide and related analogues.

When formulated as a topical emulsion, the oily phase may optionally comprise merely an emulsifier (otherwise known as an emulgent), or it may comprises a mixture of at least one emulsifier with a fat or an oil or with both a fat and an oil. Preferably, a hydrophilic emulsifier is included together with a lipophilic emulsifier which acts as a stabiliser. It is also preferred to include both an oil and a fat. Together, the emulsifier(s) with or without stabiliser(s) make up the so-called emulsifying wax, and the wax together with the oil and/or fat make up the so-called emulsifying ointment base which forms the oily dispersed phase of the cream formulations.

10

Suitable emulgents and emulsion stabilisers include Tween 60, Span 80, cetostearyl alcohol, myristyl alcohol, glyceryl monostearate and sodium lauryl sulfate. The choice of suitable oils or fats for the formulation is based on achieving the desired cosmetic properties, since the solubility of the active compound in most oils likely to be used in pharmaceutical emulsion formulations may be very low. Thus the cream should preferably be a non-greasy, non-staining and washable product with suitable consistency to avoid leakage from tubes or other containers. Straight or branched chain, mono- or dibasic alkyl esters such as di-isoadipate, isocetyl stearate, propylene glycol diester of coconut fatty acids, isopropyl myristate, decyl oleate, isopropyl palmitate, butyl stearate, 2-ethylhexyl palmitate or a blend of branched chain esters known as Crodamol CAP may be used, the last three being preferred esters. These may be used alone or in combination depending on the properties required. Alternatively, high melting point lipids such as white soft paraffin and/or liquid paraffin or other mineral oils can be used.

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Formulations suitable for rectal administration may be presented as a suppository with a suitable base comprising, for example, cocoa butter or a salicylate.

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Formulations suitable for vaginal administration may be presented as pessaries, tampons, creams, gels, pastes, foams or spray formulations containing in addition to the active ingredient, such carriers as are known in the art to be appropriate.

Formulations suitable for parenteral administration (e.g., by injection, including cutaneous, subcutaneous, intramuscular, intravenous and intradermal), include aqueous and non-aqueous isotonic, pyrogen-free, sterile injection solutions which may contain anti-oxidants, buffers, preservatives, stabilisers, bacteriostats and solutes which render the formulation isotonic with the blood of the intended recipient; and aqueous and non-aqueous sterile suspensions which may include suspending agents and thickening agents, and liposomes or other microparticulate systems which are designed to target the compound to blood components or one or more organs. Examples of suitable isotonic vehicles for use in such formulations include Sodium Chloride Injection, Ringer's Solution, or Lactated Ringer's Injection. Typically, the concentration of the active ingredient in the solution is from about 1 ng/ml to about 10 µg/ml, for example from about 10 ng/ml to about 1 µg/ml. The formulations may be presented in unit-dose or multi-dose sealed containers, for example, ampoules and vials, and may be stored in a freeze-dried (lyophilised) condition requiring only the addition of the sterile liquid carrier, for example water for injections, immediately prior to use. Extemporaneous injection solutions and suspensions may be prepared from sterile powders, granules, and tablets. Formulations may be in the form of liposomes or other microparticulate systems which are designed to target the active compound to blood components or one or more organs.

Dosage

It will be appreciated that appropriate dosages of the active compounds, and compositions comprising the active compounds, can vary from patient to patient. Determining the optimal dosage will generally involve the balancing of the level of therapeutic benefit against any risk or deleterious side effects of the treatments of the present invention. The selected dosage level will depend on a variety of factors including, but not limited to, the activity of the particular compound, the route of administration, the time of administration, the rate of excretion of the compound, the duration of the treatment, other drugs, compounds, and/or materials used in combination, and the age, sex, weight, condition, general health, and prior medical history of the patient. The amount of compound and route of

administration will ultimately be at the discretion of the physician, although generally the dosage will be to achieve local concentrations at the site of action which achieve the desired effect.

- 5 Administration *in vivo* can be effected in one dose, continuously or intermittently throughout the course of treatment. Methods of determining the most effective means and dosage of administration are well known to those of skill in the art and will vary with the formulation used for therapy, the purpose of the therapy, the target cell being treated, and the subject being treated. Single or multiple
10 administrations can be carried out with the dose level and pattern being selected by the treating physician.

- In general, a suitable dose of the active compound is in the range of about 0.1 to about 250 mg per kilogram body weight of the subject per day. Where the active
15 ingredient is a salt, an ester, prodrug, or the like, the amount administered is calculated on the basis the parent compound and so the actual weight to be used is increased proportionately.

Kits

- 20 One aspect of the invention pertains to a kit comprising (a) the active ingredient, preferably provided in a suitable container and/or with suitable packaging; and (b) instructions for use, for example, written instructions on how to administer the active compound.

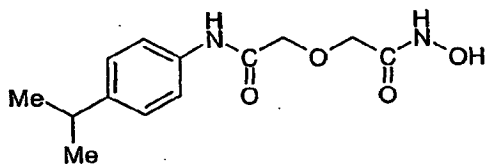
- 25 The written instructions may also include a list of indications for which the active ingredient is a suitable treatment.

EXAMPLES

- 30 The following are examples are provided solely to illustrate the present invention and are not intended to limit the scope of the invention, as described herein.

Example 1

N-Hydroxy-2-(4-isopropyl-phenylcarbamoyl)-methoxy)-acetamide
(PX083805)



- 5 ArgoGelJ-OH resin (500 mg, 0.245 mmol) was placed in a reaction vessel and was swollen by the addition of dichloromethane / pyridine (1:1, v/v) (2 ml). A solution of diglycolic anhydride (142 mg, 1.225 mmol), and 4-(dimethylamino)pyridine (DMAP) (15 mg, 0.1225 mmol) in dichloromethane / pyridine (1:1, v/v) (2 ml) was added and the resultant suspension was agitated at ambient temperature for sixteen hours. The resin was filtered and was washed with 1-methylpyrrolidine (5 ml) and alternately with methanol (4 x 5 ml) and dichloromethane (4 x 5 ml). The resin was dried and a sample was analysed by IR spectroscopy (IR(resin/cm⁻¹): 1736 (ester)).
- 15 The resin (500 mg, 0.245 mmol) obtained from the first step was placed in a reaction vessel and was swollen by the addition of 1-methylpyrrolidine (2 ml). A solution of 4-isopropylaniline (133 mg, 0.98 mmol), 1-hydroxybenzotriazole (HOBT) (66 mg, 0.49 mmol), 2-(1H-benzotriazole-1-yl)-1,1,3,3-tetramethyluronium tetrafluoroborate (TBTU) (315 mg, 0.98 mmol), *N,N*-diisopropylethylamine (DIPEA) (192 µl, 2.205 mmol) in 1-methylpyrrolidine (2 ml) was added and the resultant suspension was agitated at ambient temperature for sixteen hours. The resin was filtered and was washed with 1-methylpyrrolidine (5 ml) and alternately with methanol (4 x 5 ml) and dichloromethane (4 x 5 ml). The resin was dried and a sample was analysed by IR spectroscopy (IR(resin/cm⁻¹) 1692 (amide) and 1754 (ester)).
- 25

The resin (500 mg, 0.245 mmol) obtained from the second step was placed in a reaction vessel and was swollen by the addition of 1,4-dioxane (4 ml). A 50% wt solution of hydroxylamine in water (0.4 ml, 6.125 mmol) was added and the resultant suspension was agitated at ambient temperature for forty-eight hours.

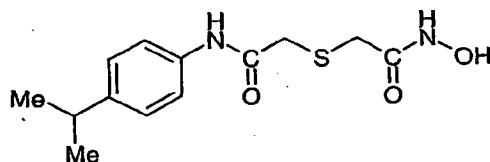
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The resin was filtered and washed with a mixture of dioxane and water (1:1, v/v) (5 ml). The filtrates were combined and the solvent was removed under reduced pressure.

- 5 The crude product obtained was purified by preparative hplc using a 150 x 21.2 mm 5 μ m Hypersil7 Elite C₁₈ column eluting with 30% ACN/70% H₂O + 0.2% TFA. The flow rate was 20 mlmin⁻¹ and the detector was set at 254 nm. The fractions that contained the desired product were concentrated under reduced pressure and the resultant residue was lyophilised from a mixture of dioxane and water to
10 afford the desired product as a white solid (5.4 mg, 8 %), t_R 3.34 (254 nm, 1.5 mlmin⁻¹, 30% ACN/70% H₂O + 0.2% TFA), m/z [ES] 265 [M - H].

Example 2

N-Hydroxy-2-(4-Isopropyl-phenylcarbamoyl)-methylsulfanyl)-acetamide
(PX089279)



- ArgoGelJ-OH resin (500 mg, 0.245 mmol) was placed in a reaction vessel and was swollen by the addition of dichloromethane / pyridine (1:1, v/v) (2 ml). A solution of thiodiglycolic anhydride (162 mg, 1.225 mmol), and 4-
20 (dimethylamino)pyridine (DMAP) (15 mg, 0.1225 mmol) in dichloromethane / pyridine (1:1, v/v) (2 ml) was added and the resultant suspension was agitated at ambient temperature for sixteen hours. The resin was filtered and was washed with 1-methylpyrrolidine (5 ml) and alternately with methanol (4 x 5 ml) and dichloromethane (4 x 5 ml). The resin was dried and a sample was analysed by
25 IR spectroscopy (IR(resin/cm⁻¹) 1732 (ester)).

- The resin (500 mg, 0.245 mmol) obtained from the first step was placed in a reaction vessel and was swollen by the addition of 1-methylpyrrolidine (2 ml). A solution of 4-isopropylaniline (133 mg, 0.98 mmol), 1-hydroxybenzotriazole
30 (HOBT) (66 mg, 0.49 mmol), 2-(1H-benzotriazole-1-yl)-1,1,3,3-tetramethyluronium

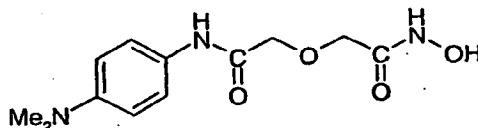
tetrafluoroborate (TBTU) (315 mg, 0.98 mmol), *N,N*-diisopropylethylamine (DIPEA) (192 μ l, 2.205 mmol) in 1-methylpyrrolidine (2 ml) was added and the resultant suspension was agitated at ambient temperature for sixteen hours. The resin was filtered and was washed with 1-methylpyrrolidine (5 ml) and alternately
5 with methanol (4 x 5 ml) and dichloromethane (4 x 5 ml). The resin was dried and a sample was analysed by IR spectroscopy (IR(resin/cm⁻¹) 1685 (amide) and 1737 (ester)).

The resin (500 mg, 0.245 mmol) obtained from the second step was placed in a
10 reaction vessel and was swollen by the addition of 1,4-dioxane (4 ml). A 50% wt solution of hydroxylamine in water (0.4 ml, 6.125 mmol) was added and the resultant suspension was agitated at ambient temperature for forty-eight hours. The resin was filtered and washed with a mixture of dioxane and water (1:1, v/v) (5 ml). The filtrates were combined and the solvent was removed under reduced
15 pressure.

The crude product obtained was purified by preparative hplc using a 150 x 21.2 mm 5 μ m Hypersil7 Elite C₁₈ column eluting with a gradient of 5% ACN/95% H₂O + 0.2% TFA to 95% ACN/5% H₂O + 0.2% TFA over 10 minutes. The flow rate
20 was 25 mlmin⁻¹ and the detector was set at 254 nm. The fractions that contained the desired product were concentrated under reduced pressure and the resultant residue was lyophilised from a mixture of dioxane and water to afford the desired product as a white solid (3.7 mg, 5 %), *t*_R 4.27 (254 nm, 3.0 mlmin⁻¹, 5% ACN/95% H₂O + 0.2% TFA to 95% ACN/5% H₂O + 0.2% TFA over 3.5 min then 2.5 min at
25 95% ACN/5% H₂O + 0.2% TFA), *m/z* [ES] 283 [M + H]⁺.

Example 3

2-(4-Dimethylamino-phenylcarbamoyl)-methoxy)-*N*-hydroxy-acetamide
(PX089281)



ArgoGelJ-OH resin (500 mg, 0.245 mmol) was placed in a reaction vessel and was swollen by the addition of dichloromethane / pyridine (1:1, v/v) (2 ml).

A solution of thiodiglycolic anhydride (162 mg, 1.225 mmol), and

4-(dimethylamino)pyridine (DMAP) (15 mg, 0.1225 mmol) in dichloromethane / pyridine (1:1, v/v) (2 ml) was added and the resultant suspension was agitated at ambient temperature for sixteen hours. The resin was filtered and was washed with 1-methylpyrrolidine (5 ml) and alternately with methanol (4 x 5 ml) and dichloromethane (4 x 5 ml). The resin was dried and a sample was analysed by IR spectroscopy (IR(resin/cm⁻¹) 1732 (ester)).

10

The resin (500 mg, 0.245 mmol) obtained from the first step was placed in a reaction vessel and was swollen by the addition of 1-methylpyrrolidine (2 ml). A solution of *N,N*-dimethylamino-1,4-phenylenediamine (133 mg, 0.98 mmol), 1-hydroxybenzotriazole (HOBT) (66 mg, 0.49 mmol), 2-(1H-benzotriazole-1-yl)-1,1,3,3-tetramethyluronium tetrafluoroborate (TBTU) (315 mg, 0.98 mmol), *N,N*-diisopropylethylamine (DIPEA) (192 µl, 2.205 mmol) in 1-methylpyrrolidine (2 ml) was added and the resultant suspension was agitated at ambient temperature for sixteen hours. The resin was filtered and was washed with 1-methylpyrrolidine (5 ml) and alternately with methanol (4 x 5 ml) and dichloromethane (4 x 5 ml). The resin was dried and a sample was analysed by IR spectroscopy (IR(resin/cm⁻¹) 1684 (amide) and 1751 (ester)).

20

The resin (500 mg, 0.245 mmol) obtained from the second step was placed in a reaction vessel and was swollen by the addition of 1,4-dioxane (4 ml). A 50% wt solution of hydroxylamine in water (0.4 ml, 6.125 mmol) was added and the resultant suspension was agitated at ambient temperature for forty-eight hours. The resin was filtered and washed with a mixture of dioxane and water (1:1, v/v) (5 ml). The filtrates were combined and the solvent was removed under reduced pressure.

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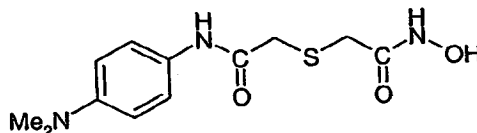
The crude product obtained was purified by preparative hplc using a 150 x 21.2 mm 5 µm Hypersil7 Elite C₁₈ column eluting with a gradient of 5% ACN/95% H₂O + 0.2% TFA to 95% ACN/5% H₂O + 0.2% TFA over 10 minutes. The flow rate

30

was 25 mlmin⁻¹ and the detector was set at 254 nm. The fractions that contained the desired product were concentrated under reduced pressure and the resultant residue was lyophilised from a mixture of dioxane and water to afford the desired product as a white solid (37.9 mg, 58 %), *t*_R 0.50 (254 nm, 3.0 mlmin⁻¹, 5% ACN/95% H₂O + 0.2% TFA to 95% ACN/5% H₂O + 0.2% TFA over 3.5 min then 2.5 min at 95% ACN/5% H₂O + 0.2% TFA).

Example 4

2-((4-Dimethylamino-phenylcarbonyl)-methylsulfanyl)-*N*-hydroxy-acetamide
(PX089278)



ArgoGelJ-OH resin (500 mg, 0.245 mmol) was placed in a reaction vessel and was swollen by the addition of dichloromethane / pyridine (1:1, v/v) (2 ml). A solution of thiodiglycolic anhydride (162 mg, 1.225 mmol), and 4-(dimethylamino)pyridine (DMAP) (15 mg, 0.1225 mmol) in dichloromethane / pyridine (1:1, v/v) (2 ml) was added and the resultant suspension was agitated at ambient temperature for sixteen hours. The resin was filtered and was washed with 1-methylpyrrolidine (5 ml) and alternately with methanol (4 x 5 ml) and dichloromethane (4 x 5 ml). The resin was dried and a sample was analysed by IR spectroscopy (IR(resin/cm⁻¹) 1732 (ester)).

The resin (500 mg, 0.245 mmol) obtained from the first step was placed in a reaction vessel and was swollen by the addition of 1-methylpyrrolidine (2 ml). A solution of *N,N*-dimethyl-1,4-phenylenediamine (133 mg, 0.98 mmol), 1-hydroxybenzotriazole (HOBT) (66 mg, 0.49 mmol), 2-(1H-benzotriazole-1-yl)-1,1,3,3-tetramethyluronium tetrafluoroborate (TBTU) (315 mg, 0.98 mmol), *N,N*-diisopropylethylamine (DIPEA) (192 µl, 2.205 mmol) in 1-methylpyrrolidine (2 ml) was added and the resultant suspension was agitated at ambient temperature for sixteen hours. The resin was filtered and was washed with 1-methylpyrrolidine (5 ml) and alternately with methanol (4 x 5 ml) and dichloromethane (4 x 5 ml). The

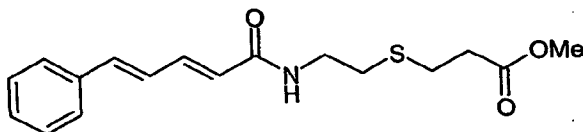
resin was dried and a sample was analysed by IR spectroscopy ($\text{IR}(\text{resin}/\text{cm}^{-1})$ 1678 (amide) and 1736 (ester)).

5 The resin (500 mg, 0.245 mmol) obtained from the second step was placed in a reaction vessel and was swollen by the addition of 1,4-dioxane (4 ml). A 50% wt solution of hydroxylamine in water (0.4 ml, 6.125 mmol) was added and the resultant suspension was agitated at ambient temperature for forty-eight hours. The resin was filtered and washed with a mixture of dioxane and water (1:1, v/v) (5 ml). The filtrates were combined and the solvent was removed under reduced
10 pressure.

The crude product obtained was purified by preparative hplc using a 150 x 21.2 mm 5 μm Hypersil7 Elite C_{18} column eluting with a gradient of 5% ACN/95% H_2O + 0.2% TFA to 95% ACN/5% H_2O + 0.2% TFA over 10 minutes. The flow rate
15 was 25 mlmin^{-1} and the detector was set at 254 nm. The fractions that contained the desired product were concentrated under reduced pressure and the resultant residue was lyophilised from a mixture of dioxane and water to afford the desired product as a brown oil (3.3 mg, 5 %), t_R 0.44 (254 nm, 3.0 mlmin^{-1} , 5% ACN/95% H_2O + 0.2% TFA to 95% ACN/5% H_2O + 0.2% TFA over 3.5 min then 2.5 min at
20 95% ACN/5% H_2O + 0.2% TFA), m/z [ES] 284 $[\text{M} + \text{H}]^+$.

Example 5

3-{2-[(2E)(4E)-5-Phenylpenta-2,4-dienoylamino]ethylsulfanyl}propionic acid methyl ester (3/28)



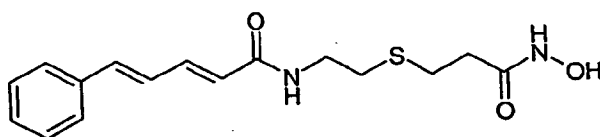
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1,1'-Carbonyldiimidazole (0.36 g, 2.2 mmol) was added to a solution of 5-phenylpenta-2E,4E-dienoic acid (1/1) (0.35 g, 2 mmol) in dry tetrahydrofuran (10 ml) and the obtained mixture was stirred for 1 hour at ambient temperature. To the mixture triethylamine (0.30 g 3.0 mmol) and methyl 3-(2-amino-ethylsulfanyl)-propionate
30 hydrochloride (2e) (2.2 mmol) were added and the resultant suspension was stirred for 6 hours at ambient temperature. The solvent was removed under

reduced pressure, to the residue water (15 ml) was added and the precipitate was filtered off, washed with water and dried, to give the title product. Yield 67%, m.p. 58-60°C. ¹H NMR (90 MHz, DMSO-d₆) δ: 2.54-2.89 (6H, m, CH₂); 2.96-3.45 (2H, m, CH₂); 3.61 (3H, s, CH₃); 6.14 (1H, d, J=14.5 Hz, CH); 6.81-7.16 (2H, m, CH-CH); 7.11-7.69 (6H, m, C₆H₅, CH); 8.18 (1H, t, J=5.4 Hz, NH).

Example 6

(2E)(4E)-5-Phenylpenta-2,4-dienoic acid [2-(2-hydroxycarbamoyl ethylsulfanyl)ethyl]amide (PX117439)



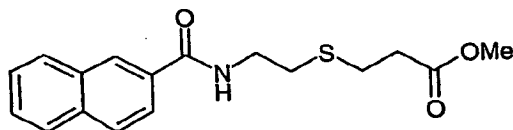
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A solution of sodium methylate (6 mmol) in methanol (5 ml) was added to a solution of hydroxylamine hydrochloride (0.28 g, 4 mmol) in methanol (8 ml). A mixture was stirred for 10 min. and the precipitate was filtered off. 3-{2-[(2E)(4E)-5-phenylpenta-2,4-dienoylamino]ethylsulfanyl}propionic acid methyl ester (3/28) (1 mmol) was added to the filtrate and the mixture was heated to the complete dissolving. The resultant mixture was stirred for 4 hours at ambient temperature and the solvent was removed under reduced pressure. The product was dissolved in water (10 ml) and acidified with 3% HCl. Precipitate was filtered off and crystallized from methanol. Yield 47%, m.p. 158-160°C. ¹H NMR (90 MHz, DMSO-d₆) δ: 2.24 (2H, t, J=6.8 Hz, CH₂); 2.56-2.87 (4H, m, CH₂); 3.29 (2H, q, J=6.0 Hz, CH₂); 6.14 (1H, d, J=15.0 Hz, CH); 6.83-7.14 (2H, m, CH-CH); 7.14-7.66 (6H, m, C₆H₅, CH); 8.21 (1H, t, J=5.5 Hz, NH); 8.74 (1H, s, NH), 10.42 (1H, s, OH). HPLC analysis on Symmetry C₁₈ column: impurities 2.6 % (column size 3.9 x 150 mm; mobile phase acetonitrile – 0.1M phosphate buffer (pH 2.5), 35:65; detector UV 230 nm; sample concentration 0.3 mg/ml). Anal. Calcd for C₁₆H₂₀N₂O₃S: C 59.98, H 6.29, N 8.74. Found: C 59.82, H 6.23, N 8.68.

20
25

Example 7

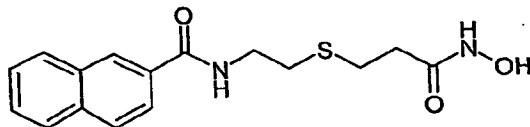
E-3-{2-[(Naphthalene-2-carbonyl)amino]ethylsulfanyl}propionic acid methyl ester
(3/29)



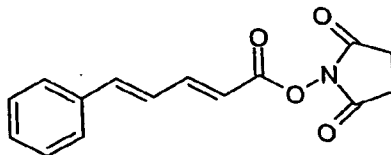
- 5 Using an analogous method, the title compound was obtained from naphthalene-2-carboxylic acid (1/29) and methyl 3-(2-amino-ethylsulfanyl)-propionate hydrochloride (2e). Yield 60%, m.p. 73-75°C. ¹H NMR (90 MHz, DMSO-d₆) δ: 2.58-2.93 (6H, m, CH₂); 3.36-3.58 (2H, m, CH₂); 3.61 (3H, s, CH₃); 7.45-7.78 (2H, m, C₁₀H₂); 7.81-8.17 (4H, m, C₁₀H₄); 8.45 (1H, s, C₁₀H); 8.78 (1H, t, J=5.2 Hz, NH).
- 10 NH).

Example 8

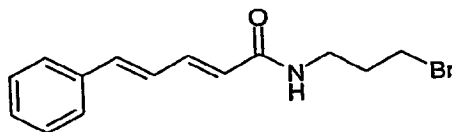
E-Naphthalene-2-carboxylic acid [2-(2-hydroxycarbamoyl)ethylsulfanyl] ethyl]amide (PX117440)



- 15 Using an analogous method, the title compound was obtained from E-3-{2-[(naphthalene-2-carbonyl)amino]ethylsulfanyl} propionic acid methyl ester (3/29). Yield 88%, m.p. 128-130°C. ¹H NMR (90 MHz, DMSO-d₆) δ: 2.27 (2H, t, J=6.7 Hz, CH₂); 2.58-2.92 (4H, m, CH₂); 3.49 (2H, q, J=5.5 Hz, CH₂); 7.52-7.74 (2H, m, C₁₀H₂); 7.87-8.18 (4H, m, C₁₀H₄); 8.47 (1H, s, C₁₀H); 8.78 (1H, t, J=5.5 Hz, NH), 10.21 (2H, br s, OH, NH). HPLC analysis on Symmetry C₁₈ column: impurities 1.0 % (column size 3.9 x 150 mm; mobile phase acetonitrile – 0.1M phosphate buffer (pH 2.5), 35:65; detector UV 230 nm; sample concentration 0.5 mg/ml). Anal. Calcd for C₁₆H₁₈N₂O₅S: C 60.36, H 5.70, N 8.80 Found: C 60.12, H 5.56, N 8.61.
- 20
- 25

Example 95-Phenyl-penta-2,4-dienoic acid 2,5-dioxo-pyrrolidin-1-yl ester (**10**)

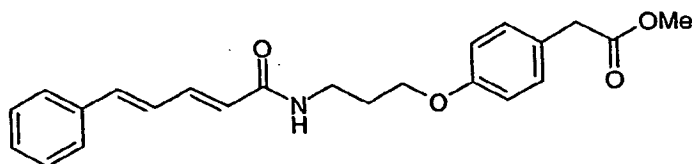
- To the solution of 5-phenyl-penta-2,4-dienoic acid (**1/1**) (8.00 g, 46.0 mmol) in anhydrous methylene chloride (350 ml) N-hydroxysuccinimide (5.29 g, 46.0 mmol) was added and the mixture was cooled in the ice bath. A solution of N,N'-dicyclohexylcarbodiimide (10.44 g, 50.6 mmol) in anhydrous methylene chloride (100 ml) slowly was added and the resulting mixture was stirred at ambient temperature for 4 hours. The precipitate was filtered off and washed with ethanol.
- The filtrate was evaporated and dried to give the title compound **10** (10.48 g, 84%). ¹H NMR (CDCl₃, HMDSO) δ: 2.85 (4H, s); 6.14 (1H, d, J=15.0 Hz); 6.75-7.18 (2H, m); 7.29-7.63 (6H, m).

Example 105-Phenyl-penta-2,4-dienoic acid (3-bromo-propyl)-amide (**11**)

- To a suspension of 3-bromopropylamine hydrobromide (1.09 g, 5.0 mmol) in anhydrous methylene chloride triethylamine (0.61 g, 6.0 mmol) was added and the mixture was stirred at ambient temperature for 30 min. The mixture was cooled in the ice bath and a solution of 5-phenyl-penta-2,4-dienoic acid 2,5-dioxo-pyrrolidin-1-yl ester (**10**) (1.35 g, 5.0 mol) in methylene chloride (20 ml) was added. The reaction mixture was stirred at ice bath temperature for 1.5 hours and at ambient temperature for 2 hours. The solvent was removed and the residue was suspended in water. The precipitate was filtered and washed with water and hexane to give the title compound **11** (1.0 g, 68%). ¹H NMR (CDCl₃, HMDSO) δ: 1.92-2.32 (2H, m); 3.31-3.67 (4H, m); 6.00 (1H, d, J=15.0 Hz); 6.07 (1H, br s); 6.78-6.96 (2H, m); 7.22-7.61 (6H, m).

Example 11

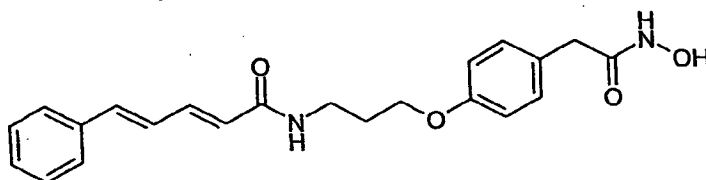
4-[3-(5-Phenyl-penta-2,4-dienoylamino)-propoxy]-phenyl}-acetic acid methyl ester
(12)



- 5 To a suspension of NaH (60% in mineral oil, 0.18 g, 4.5 mmol) in dimethylformamide (2 ml) at ice bath temperature a solution of methyl 4-hydroxyphenylacetate (0.5 g, 3.0 mmol) in dimethylformamide (2 ml) slowly was added. The resulting mixture was stirred for 1 hour at this temperature, then a solution of 5-phenylpenta-2,4-dienoic acid (3-bromo-propyl)-amide (11) was
- 10 added. The reaction mixture was stirred at 110°C for 2 hours, cooled down to room temperature and diluted with water (10 ml). The pH of the medium was brought to 5 with 1N HCl and the mixture was extracted with ethyl acetate. The organic layer was washed successively with 2N NaOH, water, brine, and dried (Na₂SO₄). The solvent was evaporated and the residue was chromatographed on
- 15 silicagel with methylene chloride-acetone (9:1) as eluent to give the title product 12 (0.240 g, 32%), m.p. 104-106°C. ¹H NMR (CDCl₃, HMDSO) δ: 1.81-2.21 (2H, m); 3.37-3.62 (4H, m); 3.66 (3H, s); 4.04 (2H, t); 5.94 (1H, d, J=15.0 Hz); 6.03 (1H, br s); 6.75-6.98 (4H, m); 7.09-7.58 (8H, m).

Example 12

(2E,4E)-N-(3-{4-[2-(Hydroxyamino)-2-oxoethyl]phenoxy}propyl)-5-phenyl-2,4-pentadienamide (PX117443)

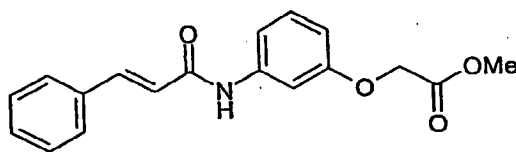


- 25 To a solution of 4-[3-(5-phenyl-penta-2,4-dienoylamino)-propoxy]-phenyl}-acetic acid methyl ester (12) (0.24 g, 0.632 mmol) in methanol (5 ml) and dioxane (2 ml) at ambient temperature a solution of hydroxylamine hydrochloride (0.176 g, 2.529 mmol) in methanol (3 ml) was added. The reaction mixture was cooled in the ice

bath and a solution of NaOH (0.20 g, 5.059 mmol) in water (1 ml) was added. The reaction mixture was stirred at ambient temperature for 1.5 hours and then pH of the medium was brought to 4 with 1N HCl. The mixture immediately was extracted with ethyl acetate (120 ml). The organic layer was washed successively with 5% NaOH, water, brine, and dried (Na₂SO₄). The solvent was partially removed, the precipitate was filtered and crystallized from acetonitrile-ethanol (1:1) to give the title compound (0.22 g, 91%). M.p. 198-201°C. ¹H NMR (DMSO-d₆, HMDSO), δ: 1.78-1.98 (2H, m, CH₂); 3.19 (2H, s, CH₂); 3.20-3.40 (2H, t, CH₂, overlapped with a signal of H₂O); 3.97 (2H, t, J=6.2 Hz, CH₂); 6.14 (1H, d, J=14.4 Hz, CH); 6.85 (2H, d, J=8.6 Hz, C₆H₂); 6.94-7.05 (2H, m, =CH-CH=); 7.09-7.42 (6H, m, C₆H₅ and -CH=); 7.50-7.61 (2H, m, C₆H₂); 8.18 (1H, t, J=5.4 Hz, NH); 8.79 (1H, s, NH); 10.59 (1H, s, OH). HPLC analysis on Symmetry C₁₈ column: impurities ~3 % (column size 3.9 x 150 mm; mobile phase acetonitrile-0.1 M phosphate buffer (40:60), pH 2.5; detector UV 230 nm; sample concentration 0.1 mg/ml, flow rate 1.0 ml/min). Anal. Calcd for C₂₂H₂₄N₂O₄: C 69.46, H 6.36, N 7.36. Found: C 69.31, H 6.34, N 7.28.

Example 13

[3-(3-Phenyl-acryloylamino)-phenoxy]-acetic acid methyl ester (8)



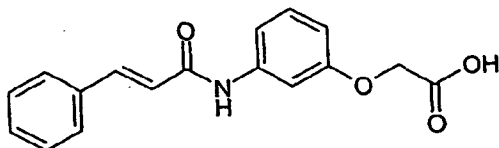
20

A mixture of methyl bromoacetate (0.230 g, 1.93 mmol), Cs₂CO₃ (0.525 g, 1.60 mmol), and N-(3-hydroxy-phenyl)-3-phenyl-acrylamide (1/91) (0.370 g, 1.55 mmol) in acetonitrile (5 ml) was stirred at 45°C for 3 hours. The reaction mixture was evaporated and the residue was partitioned between ethyl acetate and 0.1N KHSO₄. The organic layer was washed successively with water, brine, and dried (Na₂SO₄). The solvent was evaporated and the residue was chromatographed with chloroform-ethyl acetate (10:1) as eluent to give the title compound 8 (0.235 g, 48%). ¹H NMR (CDCl₃, HMDSO) δ: 3.75 (3H, s); 4.62 (2H, s); 6.53 (1H, d, J=15.6 Hz); 6.68 (1H, t, J=2.0 Hz); 7.01-7.60 (9H, m); 7.71 (1H, d, J=15.6 Hz).

30

Example 14

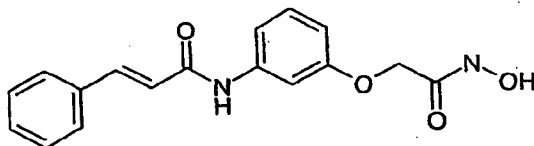
[3-(3-Phenyl-acryloylamino)-phenoxy]-acetic acid (9)



To a suspension of [3-(3-phenyl-acryloylamino)-phenoxy]-acetic acid methyl ester
5 (8) (0.345 g, 1.1 mmol) in tetrahydrofuran (6 ml) a solution of NaOH (0.130 g, 3.3
mmol) in water (6 ml) was added and the resulting mixture was stirred for 3 hours
at 35°C. Tetrahydrofuran was evaporated and the remaining water mixture was
acidified with 2N HCl to pH 3. The precipitate was filtered off and dried to give
0.300 g (92%) of the title product 9, which was used in the next step without an
10 additional purification.

Example 15

(E)-N-{3-[2-(Hydroxyamino)-2-oxoethoxy]phenyl}-3-phenyl-2-propenamide
(PX117708)



15

To a solution of [3-(3-phenyl-acryloylamino)-phenoxy]-acetic acid (9) (0.297 g, 1.0
mmol) in tetrahydrofuran (3 ml) at 0+5°C temperature (ice bath) triethylamine
(0.18 ml, 1.3 mmol) followed by iso-butyl chloroformate (0.15 ml, 1.15 mmol) were
added. The reaction mixture was stirred for 20 min. In another vessel, a mixture of
20 hydroxylamine hydrochloride (0.122 g, 1.7 mmol) and sodium methoxide (0.095
mg, 1.75 mmol) was prepared. The both mixtures were combined and the
resulting mixture was stirred at room temperature for 1 hour. The reaction mixture
was extracted with ethyl acetate (20 ml) and the organic layer was washed
successively with 1N KH₂PO₄, water, brine, and dried (Na₂SO₄). The solvent was
25 partially removed until the formation of a precipitate started. The mixture was
allowed to stand at room temperature for 6 hours. The precipitate was filtered and
washed with diethyl ether to give the title compound (0.090 g, 30%). M.p. 163°C
(dec.). ¹H NMR (DMSO-d₆, HMDSO), δ: 4.45 (s, 2H); 6.60-6.73 (m, 1H); 6.83 (d,

1H, J=15.6 Hz); 7.18-7.30 (m, 2H); 7.36-7.51 (m, 4H); 7.53-7.68 (m, 3H); 8.98 (s, 1H); 10.23 (s, 1H); 10.87 (s, 1H). HPLC analysis on Zorbax SB-C18 column: impurities ~1 % (column size 4.6 x 150 mm; mobile phase acetonitrile – 0.1% H₃PO₄, gradient from 50:50 to 100:0; detector UV 270 nm; sample concentration 0.7 mg/ml, flow rate 1.0 ml/min). Anal. Calcd for C₁₇H₁₆N₂O₄: C 65.38, H 5.16, N 8.97. Found: C 65.09, H 5.14, N 8.73.

Biological Activity

- 10 Candidate compounds were assessed for their ability to inhibit deacetylase activity (biochemical assays) and to inhibit cell proliferation (cell-based antiproliferation assays), as described below.

Primary Assay: Deacetylase Activity

- 15 Briefly, this assay relies on the release of radioactive acetate from a radioactively labelled histone fragment by the action of HDAC enzyme. Test compounds, which inhibit HDAC, reduce the yield of radioactive acetate. Signal (e.g., scintillation counts) measured in the presence and absence of a test compound
20 provide an indication of that compound's ability to inhibit HDAC activity. Decreased activity indicates increased inhibition by the test compound.

- The histone fragment was an N-terminal sequence from histone H4, and it was labelled with radioactively labelled acetyl groups using tritiated acetylcoenzyme A
25 (coA) in conjunction with an enzyme which is the histone acetyltransferase domain of the transcriptional coactivator p300. 0.33 mg of peptide H4 (the N-terminal 20 amino acids of histone H4, synthesized using conventional methods) were incubated with His6-tagged p300 histone acetyltransferase domain (amino acids 1195-1673, expressed in *E. coli* strain BLR(DE3)pLysS (Novagen, Cat. No.
30 69451-3) and 3H-acetyl coA (10 µL of 3.95 Ci/mmol; from Amersham) in a total volume of 300 µL of HAT buffer (50 mM TrisCl pH 8, 5% glycerol, 50 mM KCl, 0.1 mM ethylenediaminetetraacetic acid (EDTA), 1 mM dithiothreitol (DTT) and 1 mM 4-(2-aminoethyl)-benzenesulfonylfluoride (AEBSF)). The mixture was

incubated at 30°C for 45 min after which the His-p300 was removed using nickel-trinitriloacetic acid agarose (Qiagen, Cat No. 30210). The acetylated peptide was then separated from free acetyl coA by size exclusion chromatography on Sephadex G-15 (Sigma G-15-120), using distilled H₂O as the mobile phase.

5

After purification of the radiolabelled histone fragment, it was incubated with a source of HDAC (e.g., an extract of HeLa cells (a rich source of HDAC), recombinantly produced HDAC1 or HDAC2) and any released acetate was extracted into an organic phase and quantitatively determined using scintillation counting. By including a test compound with the source of HDAC, that compound's ability to inhibit the HDAC was determined.

10

HeLa Cell Extract

The HeLa cell extract was made from HeLa cells (ATCC Ref. No. CCL-2) by freeze-thawing three times in 60 mM TrisCl pH 8.0, 450 mM NaCl, 30% glycerol. Two cell volumes of extraction buffer were used, and particulate material was centrifuged out (20800 g, 4°C, 10 min). The supernatant extract having deacetylase activity was aliquotted and frozen for storage.

20

Recombinantly Produced HDAC1 and HDAC2

Recombinant plasmids were prepared as follows.

Full length human HDAC1 was cloned by PCR using a λ gt11 Jurkat cDNA library (Clontech-HL5012b). The amplified fragment was inserted into the EcoRI-Sall sites of pFlag-CTC vector (Sigma-E5394), in frame with the Flag tag. A second PCR was carried out in order to amplify a fragment containing the HDAC1 sequence fused to the Flag tag. The resulting fragment was subcloned into the EcoRI-Sac1 sites of the baculovirus transfer vector pAcHTL-C (Pharmingen-21466P).

30

Full length human HDAC2 was subcloned into pAcHLT-A baculovirus transfer vector (Pharmingen-21464P) by PCR amplification of the EcoRI-SacI fragment from a HDAC2-pFlag-CTC construct.

5 Recombinant protein expression and purification was performed as follows.

HDAC1 and HDAC2 recombinant baculoviruses were constructed using BaculoGold Transfection Kit (Pharmingen-554740). Transfer vectors were co-transfected into SF9 insect cells (Pharmingen-21300C). Amplification of
10 recombinant viruses was performed according to the Pharmingen Instruction Manual. SF9 cells were maintained in serum-free SF900 medium (Gibco 10902-096).

For protein production, 2×10^7 cells were infected with the appropriate recombinant
15 virus for 3 days. Cells were then harvested and spun at 3,000 rpm for 5 minutes. They were then washed twice in PBS and resuspended in 2 pellet volumes of lysis buffer (25 mM HEPES pH 7.9, 0.1 mM EDTA, 400 mM KCl, 10% glycerol, 0.1% NP-40, 1 mM AEBSF). Resuspended cells were frozen on dry ice and thawed at
20 37°C 3 times and centrifuged for 10 minutes at 14,000 rpm. The supernatant was collected and incubated with 300 μ l of 50% Ni-NTA agarose bead slurry (Qiagen-30210). Incubation was carried out at 4°C for 1 hour on a rotating wheel. The slurry was then centrifuged at 500 g for 5 minutes. Beads were washed twice in 1 ml of wash buffer (25 mM HEPES pH 7.9, 0.1 mM EDTA, 150 mM KCl, 10% glycerol, 0.1% NP-40, 1 mM AEBSF). Protein was eluted 3 times in 300 μ l elution
25 buffer (25 mM HEPES pH 7.9, 0.1 mM EDTA, 250 mM KCl, 10% glycerol, 0.1% NP-40, 1 mM AEBSF) containing increasing concentrations of imidazole: 0.2 M, 0.5 M and 1 M. Each elution was performed for 5 minutes at room temperature. Eluted protein was kept in 50% glycerol at -70°C.

30 Assay Method

A source of HDAC (e.g., 2 μ L of crude HeLa extract, 5 μ L of HDAC1 or HDAC2; in elution buffer, as above) was incubated with 3 μ L of radioactively labelled peptide

- 100 -

along with appropriate dilutions of candidate compounds (1.5 μ L) in a total volume of 150 μ L of buffer (20 mM Tris pH 7.4, 10% glycerol). The reaction was carried out at 37°C for one hour, after which the reaction was stopped by adding 20 μ L of 1 M HCl / 0.4 M sodium acetate. Then, 750 μ L of ethyl acetate was added, the samples vortexed and, after centrifugation (14000 rpm, 5 min), 600 μ L from the upper phase were transferred to a vial containing 3 mL of scintillation liquid (UltimaGold, Packard, Cat. No. 6013329). Radioactivity was measured using a Tri-Carb 2100TR Liquid Scintillation Analyzer (Packard).

10 Percent activity (% activity) for each test compound was calculated as:

$$\% \text{ activity} = \{ (S^C - B) / (S^o - B) \} \times 100$$

wherein S^C denotes signal measured in the presence of enzyme and the compound being tested, S^o denotes signal measured in the presence of enzyme but in the absence of the compound being tested, and B denotes the background signal measured in the absence of both enzyme and compound being tested. The IC50 corresponds to the concentration which achieves 50% activity.

20 IC50 data for several compounds of the present invention, as determined using this assay, are also shown in Table 1, below.

Measurement of cell viability in the presence of increasing concentration of test compound at different time points is used to assess both cytotoxicity and the effect of the compound on cell proliferation.

Secondary Assay: Cell Proliferation

30 Compounds with HDAC inhibition activity, as determined using the primary assay, were subsequently evaluated using secondary cell-based assays. The following cell lines were used:

HeLa - Human cervical adenocarcinoma cell line (ATCC ref. No. CCL-2).

K11 – HPV E7 transformed human keratinocyte line provided by Pidder Jansen-Duerr, Institut für Biomedizinische Alternsforschung, Innsbruck, Austria.

- 5 NHEK-Ad – Primary human adult keratinocyte line (Cambrex Corp., East Rutherford, NJ, USA).

JURKAT – Human T-cell line (ATCC no. TIB-152).

10 Assay Method

Cells were cultured, exposed to candidate compounds, and incubated for a time, and the number of viable cells was then assessed using the Cell Proliferation Reagent WST-1 from Boehringer Mannheim (Cat. No. 1 644 807), described
15 below.

- Cells were plated in 96-well plates at $3 \cdot 10^3$ cells/well in 100 μ L of culture medium. The following day, different concentrations of candidate compounds were added and the cells incubated at 37°C for 48 h. Subsequently, 10 μ L/well of
20 WST-1 reagent was added and the cells reincubated for 1 hour. After the incubation time, absorbance was measured.

- WST-1 is a tetrazolium salt which is cleaved to formazan dye by cellular enzymes. An expansion in the number of viable cells results in an increase in the overall
25 activity of mitochondrial dehydrogenases in the sample. This augmentation in the enzyme activity leads to an increase in the amount of formazan dye formed, which directly correlates to the number of metabolically active cells in the culture. The formazan dye produced is quantified by a scanning multiwell spectrophotometer by measuring the absorbance of the dye solution at 450 nm wavelength
30 (reference wavelength 690 nm).

Percent activity (% activity) in reducing the number of viable cells was calculated for each test compound as:

$$\% \text{ activity} = \{ (S^C - B) / (S^o - B) \} \times 100$$

5

wherein S^C denotes signal measured in the presence of the compound being tested, S^o denotes signal measured in the absence of the compound being tested, and B denotes the background signal measured in blank wells containing medium only. The IC50 corresponds to the concentration which achieves 50% activity.

10 IC50 values were calculated using the software package Prism 3.0 (GraphPad Software Inc., San Diego, CA) , setting top value at 100 and bottom value at 0.

IC50 data for several compounds of the present invention, as determined using this assay, are also shown in Table 2, below.

15

Measurement of cell viability in the presence of increasing concentration of test compound at different time points is used to assess both cytotoxicity and the effect of the compound on cell proliferation.

20 Biological Data

IC50 (or percent activity) data for several compounds of the present invention, as determined using the assays described above are summarised in Table 1 and Table 2, below.

25

Table 1 Biochemical Assay Data				
Compound		HDAC Inhibition (IC ₅₀ unless otherwise specified)		
No.	Ref.	HeLa	HDAC1	HDAC2
TSA	-	5 nM	15 nM	17 nM
1	PX083805	16% @ 1.0 μ M	-	-
2	PX089278	26% @ 500 nM	-	-
3	PX089279	55 μ M	-	-
4	PX089281	0% @ 1 μ M	-	-
5	PX117439	74% @ 500 nM	-	-
6	PX117440	43% @ 500 nM	-	-
7	PX117443	7% @ 500 nM	-	-
8	PX117708	750	-	-

Table 2 Cell-Based Antiproliferation Assay Data					
Compound		Cell Proliferation Inhibition WST-1 (IC ₅₀ unless otherwise specified)			
No.	Ref.	HeLa	K11	NHEK-AD	Jurkat
	TSA	350 nM	0.38 μ M	0.2 μ M	42 nM
	Oxamflatin	-	4.82 μ M	3.53 μ M	170 nM
	MS-275	-	9.16 μ M	3.1 μ M	365 nM
	SAHA	-	6.82 μ M	5.37 μ M	750 nM
1	PX083805	-	-	-	-
2	PX089278	-	-	-	-
3	PX089279	213 μ M	-	-	-
4	PX089281	-	-	-	-
5	PX117439	87 μ M	-	-	-
6	PX117440	-	-	-	-
7	PX117443	-	-	-	-
8	PX117708	25 μ M	1.05 μ M	-	12.7 μ M

REFERENCES

- A number of patents and publications are cited herein in order to more fully describe and disclose the invention and the state of the art to which the invention pertains. Full citations for these references are provided herein. Each of these references is incorporated herein by reference in its entirety into the present disclosure.
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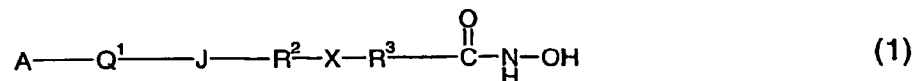
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CLAIMS

1. A compound of the formula:

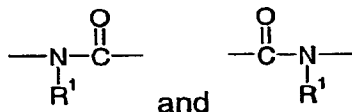


5 wherein:

A is an aryl group;

Q^1 is a covalent bond or an aryl leader group;

J is an amide linkage selected from:



10 R^1 is an amido substituent;

X is an ether heteroatom, and is -O- or -S-; and,

R^2 and R^3 are each independently an ether group;

and wherein:

15 A, is a C_{5-20} aryl group, and is optionally substituted;

the aryl leader group, if present, is a C_{1-7} alkylene group and is optionally substituted;

the amido substituent, R^1 , is hydrogen, C_{1-7} alkyl, C_{3-20} heterocyclyl, or C_{5-20} aryl;

20 each of the ether groups, R^2 and R^3 , is independently C_{1-7} alkylene; C_{5-20} arylene; C_{5-20} arylene- C_{1-7} alkylene; or C_{1-7} alkylene- C_{5-20} arylene; and is optionally substituted;

25 and pharmaceutically acceptable salts, solvates, amides, esters, ethers, chemically protected forms, and prodrugs thereof.

2. A compound according to claim 1, wherein J is -NR¹CO- and X is -O-.

3. A compound according to claim 1, wherein J is -NR¹CO- and X is -S-.

4. A compound according to claim 1, wherein Q^1 is an aryl leader group, J is $-NR^1CO-$ and X is $-O-$.
- 5 5. A compound according to claim 1, wherein Q^1 is an aryl leader group, J is $-NR^1CO-$ and X is $-S-$.
6. A compound according to claim 1, wherein Q^1 is a covalent bond, J is $-NR^1CO-$ and X is $-O-$.
- 10 7. A compound according to claim 1, wherein Q^1 is a covalent bond, J is $-NR^1CO-$ and X is $-S-$.
8. A compound according to claim 1, wherein J is $-CONR^1-$ and X is $-O-$.
- 15 9. A compound according to claim 1, wherein J is $-CONR^1-$ and X is $-S-$.
10. A compound according to claim 1, wherein Q^1 is a covalent bond, J is $-CONR^1-$ and X is $-O-$.
- 20 11. A compound according to claim 1, wherein Q^1 is a covalent bond, J is $-CONR^1-$ and X is $-S-$.
12. A compound according to claim 1, wherein Q^1 is an aryl leader group, J is $-CONR^1-$ and X is $-O-$.
- 25 13. A compound according to claim 1, wherein Q^1 is an aryl leader group, J is $-CONR^1-$ and X is $-S-$.
- 30 14. A compound according to any one of claims 1 to 13, wherein the aryl leader group, Q^1 , is a saturated C_{1-7} alkylene group.

15. A compound according to any one of claims 1 to 13, wherein the aryl leader group, Q¹, is a partially unsaturated C₁₋₇alkylene group.
- 5 16. A compound according to any one of claims 1 to 13, wherein the aryl leader group, Q¹, is an aliphatic C₁₋₇alkylene group.
17. A compound according to any one of claims 1 to 13, wherein the aryl leader group, Q¹, is a linear C₁₋₇alkylene group.
- 10 18. A compound according to any one of claims 1 to 13, wherein the aryl leader group, Q¹, is a branched C₁₋₇alkylene group.
19. A compound according to any one of claims 1 to 13, wherein the aryl leader group, Q¹, is an alicyclic C₁₋₇alkylene group.
- 15 20. A compound according to any one of claims 1 to 13, wherein the aryl leader group, Q¹, is a saturated aliphatic C₁₋₇alkylene group.
- 20 21. A compound according to any one of claims 1 to 13, wherein the aryl leader group, Q¹, is a saturated linear C₁₋₇alkylene group.
22. A compound according to any one of claims 1 to 13, wherein the aryl leader group, Q¹, is a saturated branched C₁₋₇alkylene group.
- 25 23. A compound according to any one of claims 1 to 13, wherein the aryl leader group, Q¹, is a saturated alicyclic C₁₋₇alkylene group.
24. A compound according to any one of claims 1 to 13, wherein the aryl leader group, Q¹, is a partially unsaturated aliphatic C₁₋₇alkylene group.
- 30 25. A compound according to any one of claims 1 to 13, wherein the aryl leader group, Q¹, is a partially unsaturated linear C₁₋₇alkylene group.

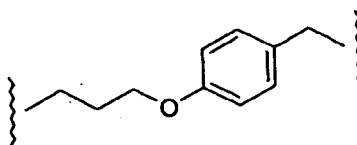
26. A compound according to any one of claims 1 to 13, wherein the aryl leader group, Q^1 , is a partially unsaturated branched C_{1-7} alkylene group.
- 5 27. A compound according to any one of claims 1 to 13, wherein the aryl leader group, Q^1 , is a partially unsaturated alicyclic C_{1-7} alkylene group.
28. A compound according to any one of claims 1 to 13, wherein the aryl leader group, Q^1 , is selected from: $-CH_2-$, $-CH=CH-$, and $-CH=CH-CH=CH-$.
- 10 29. A compound according to any one of claims 1 to 28, wherein the aryl leader group, Q^1 , is optionally substituted with one or more groups selected from: halo, hydroxy, ether, C_{5-20} aryl, acyl, amido, and oxo.
- 15 30. A compound according to any one of claims 1 to 28, wherein the aryl leader group, Q^1 , is optionally substituted with one or more groups selected from: $-F$, $-Cl$, $-Br$, $-I$, $-OH$, $-OMe$, $-OEt$, $-OPr$, $-Ph$, and $=O$.
31. A compound according to any one of claims 1 to 28, wherein the aryl leader group, Q^1 , is substituted.
- 20 32. A compound according to any one of claims 1 to 28, wherein the aryl leader group, Q^1 , is unsubstituted.
- 25 33. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently a saturated C_{1-7} alkylene group.
34. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently a partially unsaturated C_{1-7} alkylene group.
- 30 35. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently an aliphatic C_{1-7} alkylene group.

36. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently a linear C_{1-7} alkylene group.
- 5 37. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently a branched C_{1-7} alkylene group.
38. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently an alicyclic C_{1-7} alkylene group.
- 10 39. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently a saturated aliphatic C_{1-7} alkylene group.
- 15 40. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently a saturated linear C_{1-7} alkylene group.
- 20 41. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently a saturated branched C_{1-7} alkylene group.
- 25 42. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently a saturated alicyclic C_{1-7} alkylene group.
- 30 43. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently a partially unsaturated aliphatic C_{1-7} alkylene group.
44. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently a partially unsaturated linear C_{1-7} alkylene group.

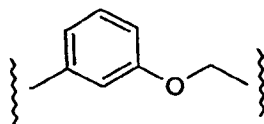
45. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently a partially unsaturated branched C_{1-7} alkylene group.
- 5 46. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently a partially unsaturated alicyclic C_{1-7} alkylene group.
- 10 47. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently $-(CH_2)_n-$, wherein n is an integer from 1 to 5.
48. A compound according to any one of claims 1 to 32, wherein the group R^2 - X - R^3 is independently selected from the following:
- 15 $-CH_2-O-CH_2-$ and $-CH_2-S-CH_2-$;
 $-CH_2-O-CH_2CH_2-$ and $-CH_2-S-CH_2CH_2-$;
 $-CH_2CH_2-O-CH_2-$ and $-CH_2CH_2-S-CH_2-$;
 $-CH_2-O-CH_2CH_2CH_2-$ and $-CH_2-S-CH_2CH_2CH_2-$;
 $-CH_2CH_2-O-CH_2CH_2-$ and $-CH_2CH_2-S-CH_2CH_2-$;
20 $-CH_2CH_2CH_2-O-CH_2-$ and $-CH_2CH_2CH_2-S-CH_2-$;
 $-CH_2-O-CH_2CH_2CH_2CH_2-$ and $-CH_2-S-CH_2CH_2CH_2CH_2-$;
 $-CH_2CH_2-O-CH_2CH_2CH_2-$ and $-CH_2CH_2-S-CH_2CH_2CH_2-$;
 $-CH_2CH_2CH_2-O-CH_2CH_2-$ and $-CH_2CH_2CH_2-S-CH_2CH_2-$;
 $-CH_2CH_2CH_2CH_2-O-CH_2-$ and $-CH_2CH_2CH_2CH_2-S-CH_2-$;
25 $-CH_2-O-CH_2CH_2CH_2CH_2CH_2-$ and $-CH_2-S-CH_2CH_2CH_2CH_2CH_2-$;
 $-CH_2CH_2-O-CH_2CH_2CH_2CH_2-$ and $-CH_2CH_2-S-CH_2CH_2CH_2CH_2-$;
 $-CH_2CH_2CH_2-O-CH_2CH_2CH_2-$ and $-CH_2CH_2CH_2-S-CH_2CH_2CH_2-$;
 $-CH_2CH_2CH_2CH_2-O-CH_2CH_2-$ and $-CH_2CH_2CH_2CH_2-S-CH_2CH_2-$;
 $-CH_2CH_2CH_2CH_2CH_2-O-CH_2-$ and $-CH_2CH_2CH_2CH_2CH_2-S-CH_2-$;
30
49. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently C_{5-20} arylene, and is optionally substituted.

50. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently C_{5-6} arylene.
- 5 51. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is phenylene.
52. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently C_{5-20} arylene- C_{1-7} alkylene or
10 C_{1-7} alkylene- C_{5-20} arylene, and is optionally substituted.
53. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently C_{5-6} arylene- C_{1-7} alkylene or
15 C_{1-7} alkylene- C_{5-6} arylene, and is optionally substituted.
54. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently C_{1-7} alkylene-phenylene.
55. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently methylene-phenylene, ethylene-
20 phenylene, propylene-phenylene, and ethenylene-phenylene.
56. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently phenylene- C_{1-7} alkylene.
- 25 57. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is independently phenylene-methylene, phenylene-ethylene, phenylene-propylene, or phenylene-ethenylene.

58. A compound according to any one of claims 1 to 32, wherein the group $-R^2$ - $X-R^3$ - is:



- 5 59. A compound according to any one of claims 1 to 32, wherein the group $-R^2$ - $X-R^3$ - is:



- 10 60. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is substituted.
61. A compound according to any one of claims 1 to 32, wherein each of the ether groups, R^2 and R^3 , is unsubstituted.
- 15 62. A compound according to any one of claims 1 to 61, wherein A is C_{5-20} heteroaryl or C_{5-20} carboaryl, and is optionally substituted.
- 20 63. A compound according to any one of claims 1 to 61, wherein A is a C_{5-20} aryl group derived from one of the following: benzene, pyridine, furan, indole, pyrrole, imidazole, naphthalene, quinoline, benzimidazole, benzothiofuran, fluorene, acridine, and carbazole.
64. A compound according to any one of claims 1 to 61, wherein A is an optionally substituted phenyl group.
- 25 65. A compound according to any one of claims 1 to 61, wherein A is a phenyl group optionally substituted with one or more of the following groups: fluoro, chloro, bromo, iodo, methyl, ethyl, isopropyl, t-butyl, cyano, trifluoromethyl, hydroxy, methoxy, ethoxy, isopropoxy, trifluoromethoxy,

phenoxy, methylthio, trifluoromethylthio, hydroxymethyl, amino, dimethylamino, diethylamino, morpholino, amido, acetamido, acetyl, nitro, sulfonamido, and phenyl.

- 5 66. A compound according to any one of claims 1 to 65, wherein the amido substituent, R^1 , is hydrogen, C_{1-7} alkyl, or C_{5-20} aryl.
67. A compound according to any one of claims 1 to 65, wherein the amido substituent, R^1 , is hydrogen or C_{1-7} alkyl.
- 10 68. A compound according to any one of claims 1 to 65, wherein the amido substituent, R^1 , is -H, -Me, or -Et.
69. A compound according to any one of claims 1 to 65, wherein the amido substituent, R^1 , is -H.
- 15 70. Compound PX083805.
71. Compound PX089278.
- 20 72. Compound PX089279.
73. Compound PX089281.
- 25 74. Compound PX117439.
75. Compound PX117440.
76. Compound PX117443.
- 30 77. Compound PX117708.

78. A composition comprising a compound according to any one of claims 1 to 77 and a pharmaceutically acceptable carrier or diluent.
- 5 79. A compound according to any one of claims 1 to 77 for use in a method of treatment of the human or animal body.
80. A compound according to any one of claims 1 to 77 for use in a method of treatment of a condition mediated by HDAC of the human or animal body.
- 10 81. A compound according to any one of claims 1 to 77 for use in a method of treatment of a proliferative condition of the human or animal body.
82. A compound according to any one of claims 1 to 77 for use in a method of treatment of cancer of the human or animal body.
- 15 83. A compound according to any one of claims 1 to 77 for use in a method of treatment of psoriasis of the human or animal body.
84. Use of a compound according to any one of claims 1 to 77 for the
20 manufacture of a medicament for use in the treatment of a condition mediated by HDAC.
85. Use of a compound according to any one of claims 1 to 77 for the
25 manufacture of a medicament for use in the treatment of a proliferative condition.
86. Use of a compound according to any one of claims 1 to 77 for the manufacture of a medicament for use in the treatment of cancer.
- 30 87. Use of a compound according to any one of claims 1 to 77 for the manufacture of a medicament for use in the treatment of psoriasis.

88. A method inhibiting HDAC in a cell comprising said cell with an effective amount of a compound according to any one of claims 1 to 77.
- 5 89. A method for the treatment of a condition mediated by HDAC comprising administering to a subject suffering from a condition mediated by HDAC a therapeutically-effective amount of a compound according to any one of claims 1 to 77.
- 10 90. A method for the treatment of a proliferative condition comprising administering to a subject suffering from a proliferative condition a therapeutically-effective amount of a compound according to any one of claims 1 to 77.
- 15 91. A method for the treatment of cancer comprising administering to a subject suffering from cancer a therapeutically-effective amount of a compound according to any one of claims 1 to 77.
- 20 92. A method for the treatment of psoriasis comprising administering to a subject suffering from psoriasis a therapeutically-effective amount of a compound according to any one of claims 1 to 77.

INTERNATIONAL SEARCH REPORT

In national Application No
PCT/GB 01/04327

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 C07C323/60 C07C259/06 A61P35/00 A61P17/06

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 C07C

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the International search (name of data base and, where practical, search terms used)
CHEM ABS Data, EPO-Internal

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	DATABASE CA 'Online! CHEMICAL ABSTRACTS SERVICE, COLUMBUS, OHIO, US; MORI, TOYOKI ET AL: "Preparation of benzothiazole derivatives as protein kinase C inhibitors" retrieved from STN Database accession no. 130:352268 XP002185425 see RN 224583-79-3 abstract & JP 11 130761 A (OHTSUKA PHARMACEUTICAL CO., LTD., JAPAN) 18 May 1999 (1999-05-18) --- -/-	1,2,6, 62, 66-69, 78,79

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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Date of the actual completion of the international search

12 December 2001

Date of mailing of the international search report

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INTERNATIONAL SEARCH REPORT

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
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INTERNATIONAL SEARCH REPORT

In Application No
PCT/GB 01/04327

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
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X	<p>FR 1 571 198 A (BOEHRINGER, C. H., SOHN) 13 June 1969 (1969-06-13)</p> <p>page 5; example 3 & CA Database accession no. 72:90063 RN 27917-61-9 BOEHRINGER, C. H., SOHN : "Antiphlogistic N-acyl-4-aminophenoxyalkylcarboxylic acid derivatives"</p> <p style="text-align: center;">--- -/-</p>	<p>1,8,10, 59, 62-69, 78,79</p>

INTERNATIONAL SEARCH REPORT

International Application No
PCT/GB 01/04327

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
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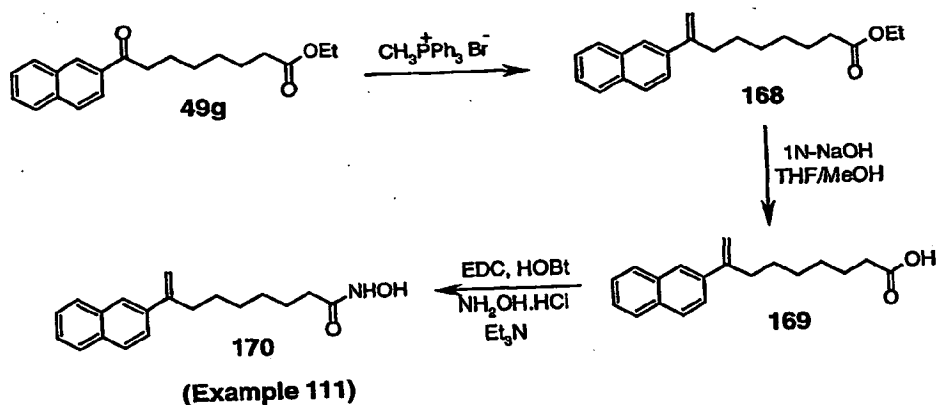
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Information on patent family members

International Application No

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**Example 111:****N-Hydroxy-8-(2-naphthyl)non-8-enamide (170)****Step 1: Ethyl 8-(2-naphthyl)non-8-enoate (168)**

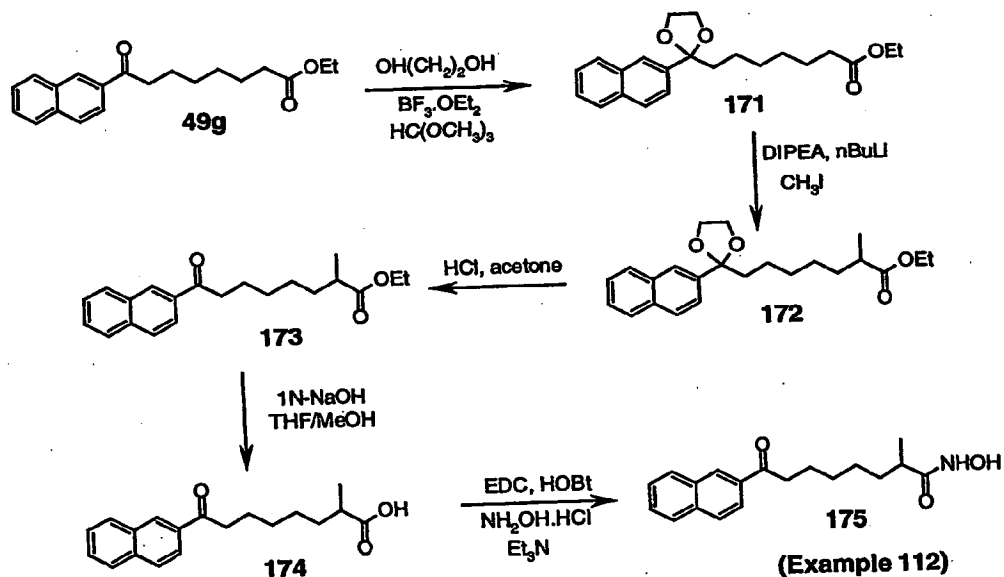
- 5 To a solution of methyltriphenylphosphonium bromide (469 mg, 1.31 mmol) in THF (6 mL) pre-cooled to 0°C under nitrogen atmosphere was added dropwise *n*-butyllithium (1.42M solution in hexanes, 905 µL, 1.28 mmol) and stirred over 20 minutes. To this a THF (4mL) solution of 49g (82 mg, 0.262 mmol) was transferred *via cannula*. The yellow mixture was stirred over 30 minutes and
- 10 quenched with a saturated aqueous solution of ammonium chloride, as the yellow color immediately disappeared. The solvent was removed by evaporation *in vacuo* and the aqueous residue was partitioned between water and ethyl acetate. The aqueous layer was then extracted with ethyl acetate and the combined organic layers were washed with brine, dried (MgSO₄) and concentrated. The
- 15 crude residue was purified by flash chromatography (5% to 8% diethyl ether in hexane) to afford the title compound 168 (43 mg) in 52% yield. ¹H NMR: (CDCl₃) δ: 7.85-7.78 (m, 4H), 7.58 (dd, J = 8.8, 1.9 Hz, 1H), 7.48-7.44 (m, 2H), 5.42 (d, J = 1.1 Hz, 1H), 5.16 (d, J = 1.4 Hz, 1H), 4.12 (q, J = 7.1 Hz, 2H), 2.62 (t, J = 6.9 Hz, 2H), 2.28 (d, J = 7.4 Hz, 2H), 1.64-1.49 (m, 4H), 1.39-1.33 (m, 4H), 1.25 (t, J = 7.1 Hz, 3H).
- 20 MS (ESI) = 311 (MH⁺).

Step 2: 8-(2-Naphthyl)non-8-enoic acid (169)

Following a procedure analogous to that described in Example 18, step 3, but substituting ester 169 for ester 49a, the title compound was obtained in 99%.
¹H NMR: (CDCl₃) δ: 7.83-7.77 (m, 4H), 7.57 (d, J = 8.8 Hz, 1H), 7.47-7.44 (m, 2H),
5.40 (s, 1H), 5.14 (s, 1H), 2.61 (t, J = 7.1 Hz, 2H), 2.33 (t, J = 7.4 Hz, 2H), 1.64-1.21
(m, 8H). MS (ESI) = 283 (MH⁺).

Step 3: N-Hydroxy-8-(2-naphthyl)non-8-enamide (170)

Following the procedure described in Example 14, step 3, but substituting
carboxylic acid 169 for 37, and using 1.1 equivalent of NH₂OH.HCl and
triethylamine each, the title compound 170 was obtained in 33% yield. ¹H NMR:
(CDCl₃) δ: 8.22 (br s, 1H), 7.80-7.77 (m, 4H), 7.56 (d, J = 8.5 Hz, 1H), 7.46-7.44 (m,
2H), 5.39 (s, 1H), 5.13 (s, 1H), 2.59 (t, J = 6.6 Hz, 2H), 2.05 (br s, 2H), 1.56-1.11 (m,
8H). HRMS: (calc.) 297.1729 (M⁺), (found) 297.1744.



Example 112:**N-Hydroxy-2-methyl-7-(2-naphthoyl)heptanamide (175)****Step 1: Ethyl-8-(2-naphthyl)-8-[2-(1,3-dioxolyl)]octanoate (171)**

To a solution of 49g (4.52 g, 14.47 mmol) in dichloromethane (140 mL) was added ethylene glycol (8.07 mL, 144.7 mmol) followed by boron trifluoride etherate (3.67 mL, 28.94 mmol). The mixture was stirred for 1 hour at room temperature. Then trimethyl orthoformate (2.73 mL, 21.71 mmol) was added and the mixture was stirred overnight. The reaction was quenched with a saturated aqueous solution of sodium bicarbonate and the layers were separated. The aqueous layer was extracted with dichloromethane and the combined organic layers were washed with brine, dried (MgSO_4) and concentrated. The crude residue was purified by flash chromatography (15% to 20% ethyl acetate/hexane), to afford the title compound 171 in 44% yield. ^1H NMR: (300 MHz, CDCl_3) δ : 7.91 (s, 1H), 7.87-7.81 (m, 3H), 7.54 (dd, $J = 8.5, 1.4$ Hz, 1H), 7.49-7.46 (m, 2H), 4.12-4.03 (m, 4H), 3.82-3.78 (m, 2H), 2.23 (t, $J = 7.4$ Hz, 2H), 1.99-1.94 (m, 2H), 1.59-1.54 (m, 2H). MS (ESI) = 357(MH^+).

Step 2: Ethyl-2-methyl-8-(2-naphthyl)-8-[2-(1,3-dioxolyl)]octanoate (172)

To a solution of freshly distilled (over sodium hydride) diisopropylamine (550 μL , 3.93 mmol) in THF (40 mL) pre-cooled to 0°C under nitrogen atmosphere was added dropwise *n*-butyllithium (1.2M solution in hexanes, 3.12 mL, 3.65 mmol). The mixture was stirred at 0°C for 20 minutes and cooled down to -78°C . Then a pre-cooled (-78°C) solution of 171 (1.0 g, 2.81 mmol) in THF (15 mL) was transferred *via cannula*. The mixture was stirred for 1 hour at -78°C . Then a pre-cooled (-78°C) solution of iodomethane (349 μL , 5.61 mmol) in THF (15 mL) was transferred *via cannula*. The resulting mixture was stirred at -78°C for 30 minutes, quenched with a saturated aqueous solution of ammonium chloride, and then warmed to room temperature. Water was added and the mixture was stirred for 4 hours, after which the THF layer had turned yellow. The mixture was concentrated *in vacuo* and the aqueous residue was partitioned between diethyl ether and water. The organic layer was washed with brine, dried (MgSO_4) and

concentrated. The crude residue was purified by flash chromatography (10% ethyl acetate in hexane) to afford the title compound 172 (950 mg) in 91% yield. ¹H NMR: (300 MHz, CDCl₃) δ: 7.91 (s, 1H), 7.87-7.81 (m, 3H), 7.54 (dd, J = 8.8, 1.9 Hz, 1H), 7.51-7.45 (m, 2H), 4.12-4.03 (m, 4H), 3.86-3.78 (m, 2H), 2.35 (sext, J = 7.1 Hz, 1H), 1.99-1.94 (m, 2H), 1.60-1.54 (m, 2H), 1.35-1.19 (m, 9H), 1.09 (d, J = 6.9 Hz, 3H). MS (ESI) = 371 (MH⁺).

Step 3 : Ethyl-2-methyl-7-(2-naphthoyl)heptanoate (173)

To a solution of 172 (950 mg, 2.56 mmol) in acetone (75 mL) was added an aqueous 1N solution of hydrochloric acid (15 mL) until a precipitate persisted, and then more acetone was added until a solution is obtained. The mixture was stirred at room temperature for 1 hour and then at 50°C for 1 hour. The mixture was concentrated *in vacuo* to a 10 mL volume. The white precipitate formed was filtered, rinsed with water to give 173. ¹H NMR: (300 MHz, CDCl₃) δ: 8.47 (s, 1H), 8.04-7.96 (m, 2H), 7.91-7.86 (m, 2H), 5.57 (quint d, J=7.7, 1.9 Hz, 2H), 4.12 (q, J = 7.1 Hz, 2H), 3.09 (t, J = 7.4 Hz, 2H), 2.46-2.39 (m, 1H), 1.82-1.51 (m, 4H), 2.45-2.34 (m, 4H), 1.24 (t, J = 7.1 Hz, 3H), 1.14 (d, J = 6.9 Hz, 3H). MS (ESI) = 327 (MH⁺).

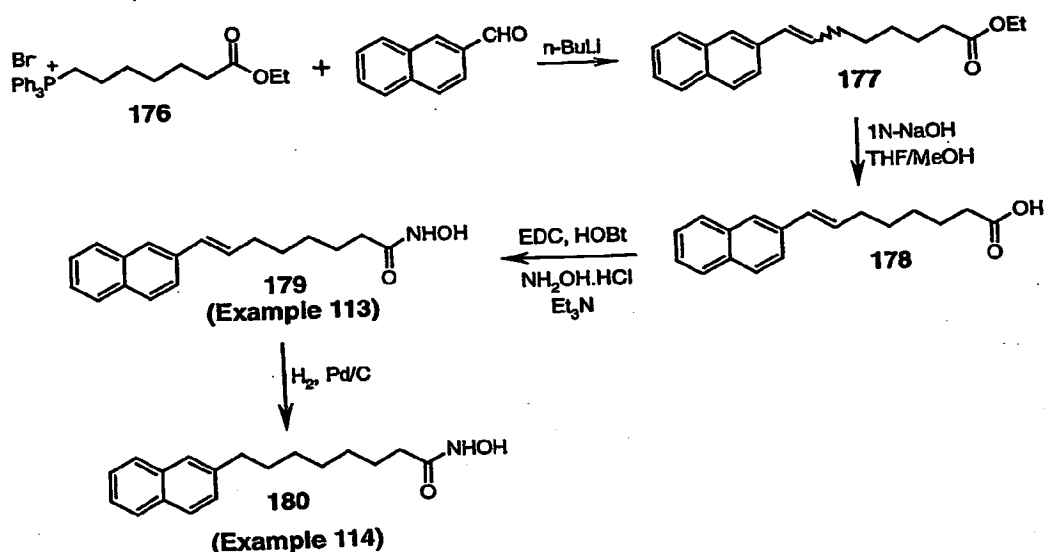
Step 4: 2-Methyl-7-(2-naphthoyl)heptanoic acid (174)

Following a procedure analogous to that described in Example 18, step 3, but substituting ester 173 for ester 49a, the title compound 174 was obtained in 73%. ¹H NMR: (300MHz, CDCl₃) δ: 8.47 (s, 1H), 8.03 (dd, J = 8.5, 1.6 Hz, 1H), 7.97 (d, J = 8.0 Hz, 1H), 7.89 (d, J = 8.5 Hz, 1H), 7.87 (d, J = 8.0 Hz, 1H), 7.57 (quint d, J = 7.7, 1.9 Hz, 2H), 3.10 (t, J = 7.1 Hz, 2H), 2.52-2.46 (m, 1H), 1.83-1.72 (m, 4H), 1.52-1.42 (m, 4H), 1.19 (d, J = 6.9 Hz, 3H). MS (ESI) = 299 (MH⁺).

Step 5 : N-Hydroxy-2-methyl-7-(2-naphthoyl)heptanamide (175)

Following a procedure analogous to that described in Example 14, step 3, but substituting carboxylic acid 174 for 37, and using 1.1 equivalent of NH₂OH.HCl and triethylamine each, the title compound 175 was obtained in 65%. ¹H NMR: (300 MHz, CDCl₃) δ: 8.47 (s, 1H), 8.02 (d, J = 8.8 Hz, 1H), 7.97 (d, J = 8.8 Hz, 1H), 7.91-7.86 (m, 2H), 7.63-7.55 (m, 2H), 3.18-3.02 (m, 2H), 2.31-2.17 (m, 1H),

1.73-1.694 (m, 2H), 1.50-1.28 (m, 4H), 1.11-1.07 (m, 3H). HRMS: (calc.) 313.1678, (found) 313.1685.



5 Example 113:

(E)-N-Hydroxy-8-(2-naphthyl)-7-octenamide (179)

Step1: (Carbethoxyhexyl)triphenylphosphonium bromide (176)

To a solution of ethyl 7-bromoheptanoate (10.0 g, 42.17 mmol) in toluene (280 mL) was added solid triphenylphosphine (11.06 g, 42.17 mmol). The solution was refluxed over 24 hours under nitrogen atmosphere and then cooled to room temperature. The supernatant was transferred into a different flask and an excess of triphenylphosphine (5.53 g, 21.09 mmol) was added. The solution was refluxed over 3 days and cooled to room temperature without stirring to favor sedimentation. The supernatant again removed by decantation and the resulting colorless oil was dried over high vacuum affording the title compound 176 (18.24 g) in 87% yield. ¹H NMR: (300 MHz, CDCl₃) δ: 7.96-7.62 (m, 9H), 7.37-7.08 (m, 6H), 4.11-4.00 (m, 2H), 3.86-3.70 (m, 2H), 2.37-2.32 (m, 4H), 2.28-2.19 (m, 2H), 1.72-1.46 (m, 6H), 1.31-1.25 (m, 2H), 1.22-1.15 (m, 3H). MS (ESI) = 419 (Phosphonium Ion).

Step2: (E/Z)-Ethyl 8-(2-naphthyl)-7-octenoate (177)

To a suspension of phosphonium salt 176 (18.14 g, 36.32 mmol) in THF (300 mL) pre-cooled to 0°C was added dropwise *n*-butyllithium (1.13M solution in hexanes, 35.4 mL, 39.95 mmol). The resulting light yellow solution was stirred at 0°C over 30 minutes and a solution of a 2-naphthaldehyde (5.67 g, 36.32 mmol) in THF (60 mL) was transferred dropwise *via cannula*. The mixture was stirred overnight, allowing to warm to room temperature. The reaction was quenched with a saturated aqueous solution of ammonium chloride. Most of the THF was removed by evaporation *in vacuo* and the aqueous residue was partitioned between diethyl ether and water. The aqueous layer was extracted with diethyl ether and the combined organic layers were successively washed with water, brine, dried over magnesium sulfate and concentrated. The crude residue was purified by flash chromatography (6% to 8% ethyl acetate in hexane) affording the title compound 177 in 34% yield as a mixture of *E* and *Z* isomers. ¹H NMR: (CDCl₃) δ: 7.83-7.75 (m, 3H), 7.71 (s, 0.5H), 7.67 (s, 0.5H), 7.57 (dd, J = 8.5, 1.9 Hz, 0.5H), 7.49-7.38 (m, 2.5H), 6.57 (d, J = 11.5 Hz, 0.5H), 6.54 (d, J = 15.9 Hz, 0.5H), 6.34 (dt, J = 15.7, 7.1 Hz, 0.5H), 5.73 (dt, J = 11.8, 7.1 Hz, 0.5H), 4.17-4.10 (m, 2H), 2.42 (qd, J = 7.4, 1.9 Hz, 1H), 2.34-2.24 (m, 3H), 1.73-1.31 (m, 6H), 1.28-1.22 (m, 3H). MS (ESI) = 297 (MH⁺).

Step3: (E)-8-(2-Naphthyl)-7-octenoic acid (178)

Following a procedure analogous to that described in Example 18, step 3, but substituting ester 177 for ester 49a, the title compound 178 was obtained in 60%. ¹H NMR: (CDCl₃) δ: 7.79-7.75 (m, 3H), 7.67 (s, 1H), 7.57 (dd, J = 8.5, 1.7 Hz, 1H), 7.47-7.38 (m, 2H), 6.54 (d, J = 15.7 Hz, 1H), 6.34 (dt, J = 9.0, 6.6 Hz, 1H), 2.38 (t, J = 7.4 Hz, 2H), 2.28 (q, J = 6.6 Hz, 1H), 1.74-1.64 (m, 2H), 1.57-1.40 (m, 4H). MS (ESI) = 269 (MH⁺).

Step 4: (E)-N-Hydroxy-8-(2-naphthyl)-7-octenamide (179)

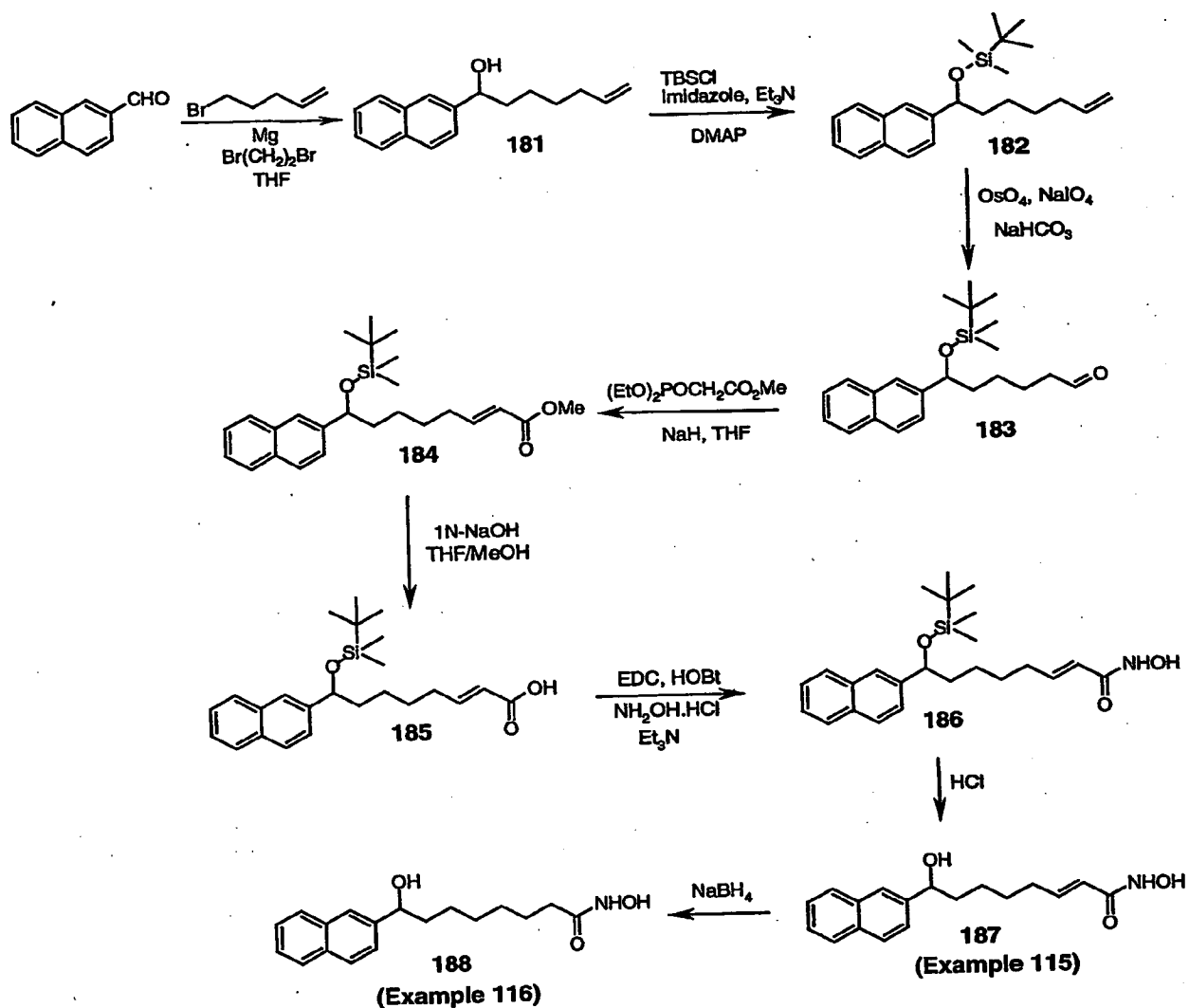
Following a procedure analogous to that described in Example 14, step 3, but substituting carboxylic acid 178 for 37, and using 1.1 equivalent of

NH₂OH.HCl and triethylamine each, the title compound 179 was obtained in 37% yield. ¹H NMR: (CD₃OD) δ: 7.79-7.74 (m, 3H), 7.67 (s, 1H), 7.60 (d, J = 8.8 Hz, 1H), 7.45-7.36 (m, 2H), 6.56 (d, J = 15.7 Hz, 1H), 6.38 (dt, J = 15.7, 6.9 Hz, 1H), 2.28 (q, J = 6.3 Hz, 2H), 2.11 (t, J = 7.4 Hz, 2H), 1.72-1.62 (m, 2H), 1.58-1.50 (m, 2H), 1.46-1.40 (m, 2H). HRMS (calc.): 283.1572, (found): 283.1567.

Example 114:

N-Hydroxy-8-(2-naphthyl)octanamide (180)

To a solution of compound 179 (30 mg, 0.106 mmol) in methanol (2 mL) was added a catalytic amount of 10% palladium on charcoal. The mixture was degassed and applied under hydrogen atmosphere and stirred for 15 minutes. Hydrogen was evacuated by vacuum and the mixture was filtered through Celite and rinsed with methanol. The filtrate was concentrated *in vacuo*, affording the title compound 180 (29.1 mg) in 97% yield. ¹H NMR: (300 MHz, CD₃OD) δ: 7.80-7.74 (m, 3H), 7.60 (s, 1H), 7.44-7.37 (m, 2H), 7.33 (dd, J = 8.2, 1.6 Hz, 1H), 2.77 (t, J = 7.4 Hz, 2H), 2.07 (t, J = 7.1 Hz, 2H), 1.71-1.64 (m, 2H), 1.62-1.57 (m, 2H), 1.37-1.28 (m, 6H). HRMS (calc.): 285.1729, (found): 285.1727.

**Example 115:****N-Hydroxy-8-hydroxy-8-(2-naphthyl)-2-octenamide (187)****5 Step1: 1-(2-Naphthyl)-6-heptenol (181)**

A flame-dried round-bottomed flask was charged with magnesium turnings (1.23 g, 50.75 mmol) and stirred under vacuum for 15 minutes and then applied under nitrogen atmosphere. THF (70 mL) was added followed by 1,2-dibromoethane (381 mg, 2.03 mmol), and the mixture was brought to the boiling point for 1 minute, then and cooled down to room temperature. A solution of 6-

10

bromohexene (8.28 g, 50.75 mmol) in THF (30 mL) was transferred *via cannula* to the magnesium flask and the mixture was refluxed overnight and cooled to -78°C . A solution of 2-naphthaldehyde (6.10 g, 39.04 mmol) in THF (30 mL) was transferred *via cannula* to the Grignard reagent and the mixture was slowly warmed to 0°C over 3 hours and kept at that temperature for 2 hours. The reaction was quenched with a saturated aqueous ammonium chloride solution and THF was evaporated *in vacuo*. The aqueous residue was poured into a separating funnel containing water and the compound was extracted with diethyl ether (2 x 100 mL). The combined organic layers were washed with brine, dried over magnesium sulfate and concentrated. The residue was purified by flash chromatography (10% to 15% ethyl acetate in hexane), affording the title compound 181 in 89% yield (8.33g). ^1H NMR: (300 MHz, CDCl_3) δ : 7.85-7.78 (m, 4H), 7.51-7.44 (m, 3H), 5.85-5.72 (m, 1H), 5.02-4.91 (m, 2H), 4.87-4.82 (m, 2H), 2.08-2.01 (m, 2H), 1.94-1.80 (m, 2H), 1.53-1.23 (m, 4H). MS (ESI) = 223 ($\text{MH}^+ - \text{H}_2\text{O}$).

15 Step2: O-(*t*-Butyldimethylsilyl)-1-(2-naphthyl)-6-heptenol (182)

To a solution of alcohol 181 (1.0g, 4.16 mmol) in dichloromethane (40mL) at room temperature under nitrogen atmosphere were successively added solid *t*-butyldimethylsilyl chloride (815 mg, 5.41 mmol), imidazole (340 mg, 4.99 mmol), and a catalytic amount of 2,6-dimethylaminopyridine. The mixture was stirred over 1 hour and triethylamine (696 μL , 4.99 mmol) was added. The mixture was stirred for 2 days. The reaction was quenched with water and the layers separated. The aqueous layer was extracted with dichloromethane and the combined organic layers were washed successively with a 1N HCl (2 x 10 mL), a saturated solution of sodium bicarbonate and brine, dried (MgSO_4) and concentrated *in vacuo*. The crude residue was purified by flash chromatography using hexane as the eluent, and affording the title compound 182 in 84% yield (1.24 g). ^1H NMR: (300 MHz, CDCl_3) δ : 7.83-7.79 (m, 3H), 7.70 (s, 1H), 7.47-7.40 (m, 3H), 5.78 (ddt, $J = 17.0, 10.4, 6.6$ Hz, 1H), 5.85-5.71 (m, 2H), 4.79 (dd, $J = 6.9, 4.9$ Hz, 1H), 2.04-1.98 (m, 2H), 1.80-1.54 (m, 2H), 0.90 (s, 9H), 0.04 (s, 3H), -0.14 (s, 3H). MS (ESI) = 223 ($\text{MH}^+ - \text{TBSO}$).

Step3: 6-t-Butyldimethylsilyloxy-6-(2-naphthyl)hexanal (183)

To a solution of alkene 182 (1.05 g, 3.23 mmol) in *t*-butyl alcohol (50mL) were successively added water (10 mL), solid sodium bicarbonate (2.71 g, 32.3 mmol), sodium periodate (4.15 g, 19.38 mmol), and osmium tetroxide (8 mg, 0.032 mmol). The mixture turned light brown and a precipitate formed in large amount, making magnetic stirring impossible so the mixture was shaken by hand every 10 minutes for 3.5 hours and the brownish color almost disappeared. The reaction was quenched with a 10% aqueous solution of sodium thiosulfate. After 30 minutes, the white precipitate was filtered off and rinsed with diethyl ether. The filtrate was concentrated *in vacuo*. The residue was partitioned between diethyl ether and water. The aqueous layer was extracted (2 x 50 mL) with diethyl ether. The combined organic layers were washed successively with a 10% aqueous solution of sodium thiosulfate, water and brine, dried (MgSO₄), and concentrated *in vacuo*, yielding the title compound 183 as a crude oil in 91% yield (1.05 g). ¹H NMR: (300 MHz, CDCl₃) δ: 9.73 (t, J = 1.9 Hz, 1H), 7.83-7.78 (m, 3H), 7.69 (s, 1H), 7.49-7.43 (m, 3H), 4.80 (t, J = 6.9 Hz, 1H), 2.39 (td, J = 7.1, 1.6 Hz, 2H), 1.81-1.55 (m, 4H), 1.43-1.25 (m, 2H) 0.90 (s, 9H), 0.04 (s, 3H), -0.15 (s, 3H). MS (ESI) 225 (M+1-TBSO).

Step 4: Methyl 8-t-butyldimethylsilyloxy-8-(2-naphthyl)-2-octenoate (184)

To a suspension of sodium hydride (60% in mineral oil, 177 mg, 4.42 mmol) in THF (25 mL) at room temperature under nitrogen atmosphere was added dropwise neat methyl diethylphosphonoacetate (811 µL, 4.42 mmol). The resulting solution was stirred for 10 minutes while cooling down to 0°C. A solution of crude aldehyde 183 (1.05 g, 2.94 mmol) in THF (10 mL) was transferred *via cannula*. The resulting solution was stirred for 15 minutes at 0°C and quenched with a saturated aqueous solution of ammonium chloride. The THF was evaporated *in vacuo* and the aqueous residue was partitioned between diethyl ether and water. The aqueous layer was extracted with diethyl ether and the combined organic layers were washed with brine, dried (MgSO₄) and concentrated *in vacuo*. The crude residue was purified by flash chromatography

(5% ethyl acetate/hexane), affording the title compound 184 in 63% yield (1.30 g).
1H NMR: (300 MHz, CDCl₃) δ : 7.83-7.78 (m, 3H), 7.69 (s, 1H), 7.49-7.41 (m, 3H),
6.94 (dt, J = 15.4, 7.1 Hz, 1H), 5.79 (dt, J = 15.7, 1.1 Hz, 1H), 4.79 (dd, J = 7.1, 5.2 Hz,
1H), 3.71 (s, 3H), 2.16 (q, J = 7.4 Hz, 2H), 1.81-1.62 (m, 2H), 1.48-1.26 (m, 4H), 0.90
5 (s, 9H), 0.04 (s, 3H), -0.15 (s, 3H). MS (ESI) = 413 (MH⁺).

Step 5: 8-t-Butyldimethylsilyloxy-8-(2-naphthyl)-2-octenoic acid (185)

Following a procedure analogous to that described in Example 18, step 3,
but substituting ester 184 for ester 49a, the title compound 185 was obtained in
100%. 1H NMR: (CDCl₃) δ : 7.83-7.78 (m, 3H), 7.72 (s, 1H), 7.48-7.40 (m, 3H), 6.74
10 (dt, J = 15.4, 7.1 Hz, 1H), 5.78 (d, J = 15.4 Hz, 1H), 4.88-4.84 (m, 1H), 2.17-2.11 (m,
2H), 1.82-1.69 (m, 2H), 1.47-1.28 (m, 4H), 0.89 (s, 9H), 0.06 (s, 3H), -0.16 (s, 3H). MS
(ESI) = 421 (M⁺+Na).

Step 6: N-Hydroxy-8-t-butyldimethylsilyloxy-8-(2-naphthyl)-2-octenamide (186)

Following a procedure analogous to that described in Example 14, step 3,
15 but substituting carboxylic acid 185 for 37, and using 1.1 equivalent of
NH₂OH.HCl and triethylamine each, the title compound 186 was obtained in 44%
yield. 1H NMR: (CD₃OD) δ : 7.83-7.72 (m, 4H), 7.48-7.40 (m, 3H), 6.77 (dt, J = 15.1,
7.1 Hz, 1H), 5.75 (d, J = 15.4 Hz, 1H), 4.91-4.79 (m, 1H), 2.18-2.14 (m, 2H), 1.79-1.68
(m, 2H), 1.48-1.39 (m, 2H), 1.28-1.21 (m, 2H), 0.89 (s, 9H), 0.06 (s, 3H), -0.16 (s, 3H).
20 MS (ESI) = 436 (M⁺+Na).

Step 7: N-Hydroxy-8-hydroxy-8-(2-naphthyl)-2-octenamide (187)

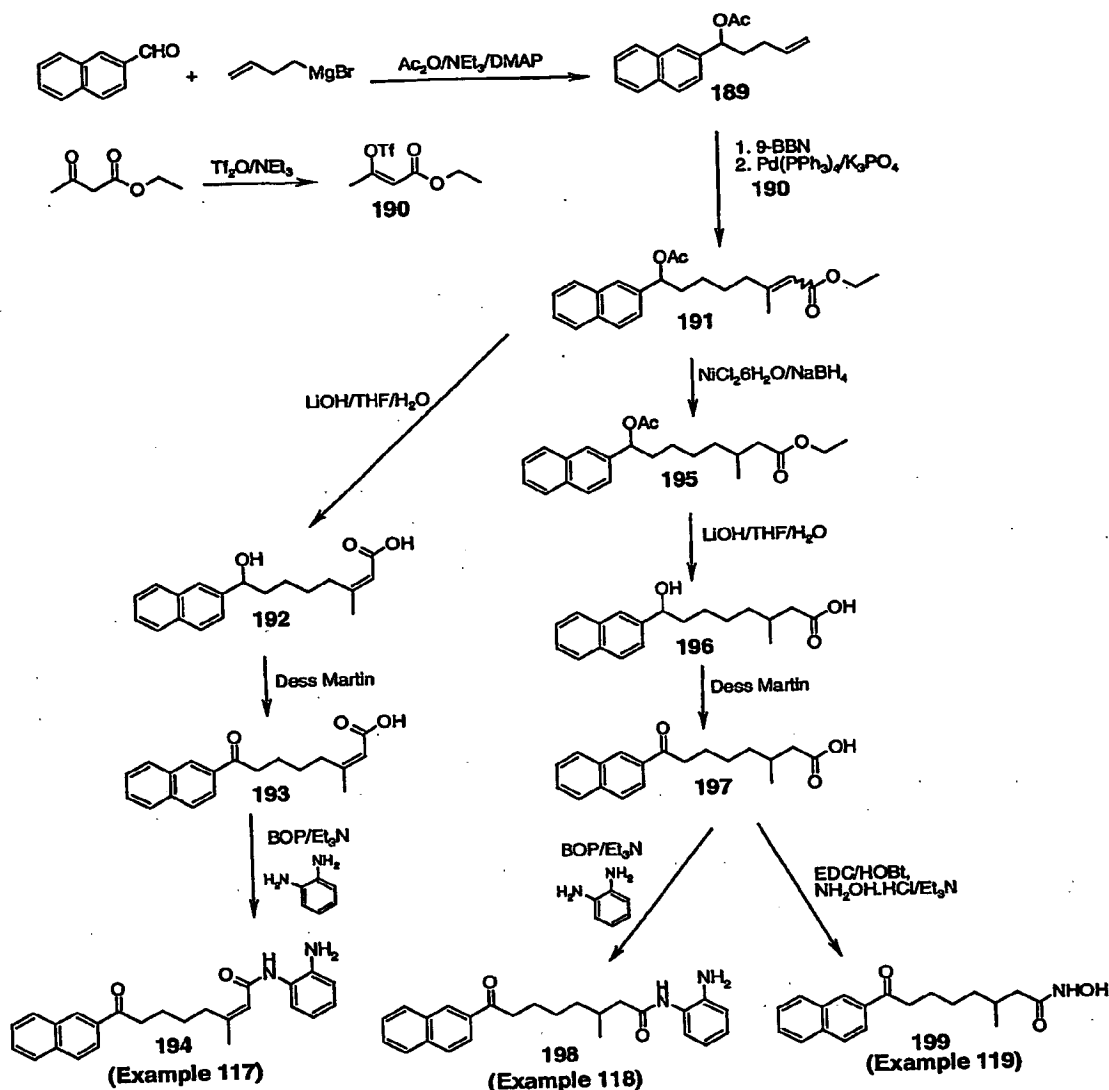
To a solution of the silyl ether 186 (300 mg, 0.725 mmol) in THF (3mL) at
room temperature was added 1N aqueous solution of hydrochloric acid (3mL).
The mixture was stirred for 3 hours at 30°C and then added a large volume of
25 ethyl acetate and water. The water layer was washed with ethyl acetate and the
combined organic layers were washed with brine, dried (MgSO₄) and
concentrated *in vacuo*. The crude residue was purified by flash chromatography
(75% to 100% ethyl acetate/hexane) affording unsatisfactory purity. So the
compound was purified again using 8% methanol in dichloromethane affording

the title compound 187 in 33% yield (77 mg). ¹H NMR: (300 MHz, CD₃OD) δ: 7.87-7.71 (m, 4H), 7.52-7.40 (m, 3H), 6.77 (dt, J = 15.1, 6.9 Hz, 1H), 5.75 (d, J = 15.1 Hz, 1H), 4.76 (t, J = 6.9 Hz, 1H), 2.17-2.10 (m, 2H), 1.84-1.77 (m, 2H), 1.49-1.27 (m, 4H), 1.88-1.75 (m, 4H). MS (ESI) = 282 (M⁺-OH), 322 (M⁺+Na).

5 **Example 116:**

N-Hydroxy-8-hydroxy-8-(2-naphthyl)octanamide (188)

To a solution of 187 (20 mg, 0.067 mmol) in methanol (670 μL) was added solid sodium borohydride (2.5 mg, 0.067 mmol). The mixture was stirred for 15 minutes and the solvent was evaporated *in vacuo*. The residue was partitioned between ethyl acetate and water. The organic layer was dried (MgSO₄) and concentrated. Purification by flash silica gel chromatography (6% methanol in dichloromethane) afforded the title compound 188 in 46% yield (9.1 mg). ¹H NMR: (300 MHz, CD₃OD) δ: 7.84-7.76 (m, 4H), 7.50-7.40 (m, 3H), 4.75 (t, J = 6.6 Hz, 1H), 2.05 (t, J = 7.4 Hz, 2H), 1.86-1.76 (m, 2H), 1.62-1.55 (m, 2H), 1.43-1.23 (m, 6H).
15 HRMS: (calc.) 283.1572 (M-H₂O), (found) 283.1581.

**Example 117:**

(Z)-N-(2-Aminophenyl)-3-methyl-7-(2-naphthoyl)-2-heptenamide (194)

Step 1: 1-O-Acetyl-2-naphthyl-4-pentenol (189)

- 5 Following a procedure analogous to that described in Example 115, step 1, but substituting 4-bromobutene for 6-bromohexene, the resulting alcohol (2.16g, 10.16 mmol) was dissolved in dichloromethane (80 ml) and was added Et₃N (5.7 ml, 40.6 mmol), acetic anhydride (2.9 ml, 30.5 mmol) and DMAP (62 mg, 5%) at 0°C. The solution was warmed up at room temperature and stirred for 5 hours.
- 10 The resulting solution was washed with a saturated solution of NaHCO₃ (2 x 50

mL), brine and then dried over Na_2SO_4 and concentrated to give the product 189 (2.6g, 99%). ^{13}C NMR: (75 MHz, CDCl_3) δ : 21.2, 29.6, 35.2, 75.5, 115.2, 124.2, 125.7, 125.9, 126.1, 127.6, 127.9, 128.2, 133.0, 133.0, 137.3, 137.7, 170.2. MS (ESI) = 277 ($\text{M}^+ + \text{Na}$).

5 Step 2: (E/Z)-Ethyl-3-trifluoromethanesulfonyloxy-2-butenate (190)

To a solution of ethyl acetoacetate (545 μl , 5mmol) in dichloromethane at -78°C was added Et_3N (2.1 ml, 15 mmol). After for 2 hours Tf_2O (1.1 ml, 3.9 mmol) was added and the solution was stirred at -78°C for 5 hours. The solution was warmed up at room temperature and was diluted with anhydrous methanol. The oil was filtered through a pad of silica gel eluting with ether and concentrated to give yellow oil. Purification by column chromatography over silica gel gave the following compounds 190: E-isomer (575 mg, 46%), Z-isomer (409 mg, 33%). ^{13}C NMR: (75 MHz, CD_3OD) δ : 25.2, 25.3, 28.7, 33.7, 39.1, 117.4, 124.7, 127.8, 128.7, 129.0, 129.5, 130.6, 131.0, 137.8, 161.9, 170.1. MS (ESI) = 297 (MH^+).

15 Step 3: (E/Z)-Ethyl-3-methyl-8-acetoxy-8-(2-naphthyl)-2-octenoate (191)

An oven-dried flask equipped with a reflux condenser and a septum inlet was flushed with nitrogen and charged with a solution of 9-BBN (0.5M, 0.66 mmol) and then alkene 189 (168 mg, 0.66 mmol) at 0°C . The mixture was warmed up slowly to room temperature and stirred for 6 hours. To this solution were added dioxane (4 ml), powder K_3PO_4 (191 mg, 0.90 mmol), $\text{Pd}(\text{PPh}_3)_4$ (18 mg, 0.015 mmol), and triflate 190 (solution in dioxane 0.18M, 3.42 ml, 0.60 mmol). The mixture was heated at 85°C for 17 hours. Then the mixture was diluted with hexane at room temperature, and the residual borane was oxidized with 3 M NaOH (1 ml) and 30% H_2O_2 (1 ml) for 1 hour. The product was extracted, washed with brine, dried over Na_2SO_4 and finally isolated by chromatography over silica gel (hexanes: Et_2O : 10:1 to 1:1) to give 70% as mixture of E and Z-isomers 191 in a 1:1.1 ratio. The 20% of Z-isomer was isolated from the mixture: ^{13}C NMR: (75 MHz, CDCl_3) δ : 21.1, 24.9, 25.5, 27.6, 32.9, 35.8, 50.6, 75.9, 115.6, 124.1, 125.5, 125.8,

126.0, 127.5, 127.8, 128.1, 132.8, 132.9, 137.9, 160.4, 166.5, 170.2. MS (ESI) = 375 (M⁺+Na).

Step 4: (Z)-8-Hydroxy-3-methyl-8-(2-naphthyl)-2-octenoic acid (192)

Following a procedure described in Example 1, step 4, but substituting
5 ester **191** for ester **3**, the compound **192** was obtained in 99% yield. ¹³C NMR: (75 MHz, CDCl₃) δ 25.3, 26.9, 29.1, 34.0, 39.7, 75.1, 117.1, 125.3, 125.6, 126.5, 126.9, 128.5, 128.8, 128.9, 134.3, 134.7, 143.9, 162.2, 169.7. MS (ESI) = 281 (MH⁺-H₂O)

Step 5: (Z)-3-Methyl-7-(2-naphthoyl)-2-heptenoic acid (193)

To a solution of **192** (98 mg, 0.33 mmol) in dry dichloromethane (6 mL) was
10 added a solution of Dess-Martin reagent (162 mg, 0.38 mmol) in dry dichloromethane (1 mL) *via cannula*. The mixture was stirred at room temperature for 24 hours. The mixture was diluted with ether and NaHSO₃ (221 mg) in aqueous solution was added with stirring until the phases are homogeneous. The product was extracted, washed with 5% KHSO₄, brine, dried over Na₂SO₄, and
15 finally isolated by chromatography over silica gel (hexanes:Et₂O: 4:1 to 1:1) to give **193** in 94% yield. ¹³CNMR: (75 MHz, CD₃OD) δ: 25.2, 25.3, 28.7, 33.7, 39.1, 117.4, 124.7, 127.8, 128.7, 129.0, 129.5, 130.6, 131.0, 137.8, 161.9, 170.1. MS (ESI) = 297 (MH⁺).

Step 6: (Z)-N-(2-Aminophenyl)-3-methyl-7-naphthoyl-2-heptenamide (194)

Following a procedure described in Example 35, but substituting
20 respectively acid **193** for acid **80** and 1,2-phenylenediamine for aniline, the compound **194** was obtained in 99% yield. ¹H NMR: (300 MHz, CD₃OD): 8.01 (d, 2H, J = 8.0 Hz), 7.92 (d, 2H, J = 8.8 Hz), 7.65-7.54 (m, 3H), 7.09 (d, J=8.0 Hz, 1H), 6.68 (dd, J = 7.7, 7.4 Hz, 1H), 5.97 (s, 1H), 3.22 (t, J = 7.1 Hz, 2H), 2.82 (dd, J = 7.4, 7.7 Hz, 2H), 1.91-1.80 (m, 2H), 1.77-1.63 (m, 2H). ¹³C NMR: (75 MHz, CH₃OD): 25.1,
25 25.2, 28.7, 33.4, 39.1, 118.6, 119.6, 124.7, 125.4, 126.9, 127.8, 127.9, 128.7, 129.4, 129.5, 130.7, 131.1, 133.9, 135.5, 136.9, 140.8, 143.0, 157.6, 167.5, 202.6

Example 118:**(Z)-N-(2-Aminophenyl)-3-methyl-7-(2-naphthoyl)heptanamide (198)****Step 1: Ethyl-3-methyl-8-O-acetoxy-8-(2-naphthyl)-2-octenoate (195)**

To a cold (ice bath) solution of 191 (126 mg, 0.36 mmol) in dry CH_3OH (4 ml) were added 3 ml of a 4% CH_3OH solution of $\text{NiCl}_2 \cdot 6\text{H}_2\text{O}$ (0.6 mmol). The solution was stirred for 30 min at 0°C and then treated portionwise with NaBH_4 (46 mg, 1.22 mmol). The resulting black solution was stirred for additional 1 hour at 0°C , and then the ice bath was removed. After 18 hours at room temperature the mixture was treated with NaHCO_3 , filtered through Celite, and concentrated. The residue was partitioned between saturated NaHCO_3 solution and dichloromethane. The combined organic layers were dried Na_2SO_4 and concentrated to give crude 195. Flash chromatography on silica gel of the residue afforded 67% of 195 (86 mg). ^{13}C NMR: (75 MHz, CDCl_3) δ : 19.5, 25.5, 26.5, 30.1, 36.0, 36.4, 41.1, 41.4, 51.3, 76.1, 124.2, 125.6, 125.9, 127.6, 127.9, 128.2, 132.9, 133.0, 137.9, 170.3, 173.6. MS (ESI) = 379 ($\text{M}^+ + \text{Na}$).

Step 2: 8-Hydroxy-3-methyl-8-(2-naphthyl)-octanoic acid (196)

Following a procedure described in Example 1, step 4, but substituting ester 195 for ester 3, the compound 196 was obtained in 99% yield. ^{13}C NMR: (75 MHz, CDCl_3) δ : 20.0, 27.0, 27.9, 31.4, 37.6, 39.9, 42.6, 75.2, 125.3, 125.6, 126.5, 126.9, 128.6, 128.8, 128.9, 134.3, 134.7, 143.9, 177.2. MS (ESI) = 323 ($\text{M}^+ + \text{Na}^+$).

Step 3: 3-Methyl-7-naphthoyl-heptanoic acid (197)

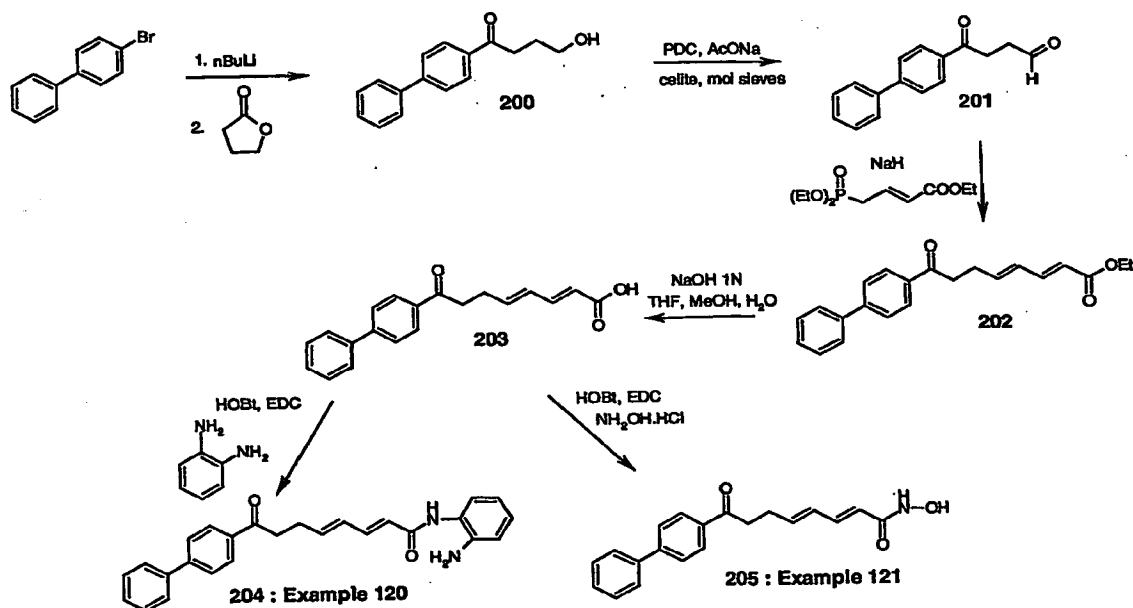
Following a procedure described in Example 117, step 5, but substituting 196 for 192, the compound 197 was obtained in 94% yield. ^1H NMR: (300 MHz, CDCl_3) δ 8.06-7.55 (m, 7H), 3.32 (br s, 1H), 3.16 (t, $J = 6.9\text{ Hz}$, 1H), 2.32 (dd, $J = 6.0, 14.0\text{ Hz}$, 1H), 2.14-2.06 (m, 1H), 1.96-1.95 (m, 1H), 1.78-1.73 (m, 2H), 1.47-1.29 (m, 3H), 0.97 (d, $J = 6.04\text{ Hz}$, 3H). MS (ESI) = 299 (MH^+).

Step 4: N-(2-Aminophenyl)-3-Methyl-7-naphthoyl-heptanamide (198)

Following a procedure described in Example 35, but substituting acid 193 for acid 80 1,2-phenylenediamine for aniline, the compound 198 was obtained in 87% yield. ¹H NMR: (300 MHz, CD₃OD): 8.05-7.91 (m, 4H), 7.67-7.53 (m, 2H), 7.10-6.68 (m, 5H), 3.33-3.31 (m, 2H), 3.19 (dd, J = 6.9, 7.4 Hz, 2H), 2.47-1.29 (m, 7H), 1.05 (d, J = 6.6 Hz, 3H), ¹³C NMR (75 MHz, CD₃OD): 20.1, 25.8, 27.7, 32.1, 37.7, 39.4, 44.87, 118.6, 119.5, 124.7, 127.1, 127.8, 128.3, 128.8, 129.5, 129.6, 130.7, 131.1, 134.1, 137.1, 174.6, 202.7.

Example 119:**10 N-Hydroxy-3-methyl-7-(2-naphthoyl)heptanamide (199)**

Following a procedure analogous to that described in Example 14, step 3, but substituting carboxylic acid 197 for 37, and using 1.1 equivalent of NH₂OH.HCl and triethylamine each, the title compound 199 was obtained in 63% yield. ¹H NMR: (300 MHz, CD₃OD): 8.07-7.58 (m, 7H), 3.32 (t, J = 1.7 Hz, 2H), 3.18 (dd, J = 7.4, 6.7 Hz, 2H), 2.13-1.30 (m, 7H), 0.95 (d, J = 6.3 Hz, 3H). ¹³C NMR (75 MHz, CD₃OD): 19.7, 25.7, 27.6, 31.7, 37.6, 39.4, 41.4, 124.7, 127.9, 128.7, 129.4, 129.6, 130.7, 131.1, 134.1, 135.6, 137.1, 172.4, 202.8.



Example 120:**(E,E)-N-(2-Aminophenyl)-7-[(4-biphenyl)carbonyl]-2,4-heptadienamide (204)****Step 1: 1-(4-Biphenyl)-4-hydroxybutanone (200)**

To a solution of bromobiphenyl (1 g, 4.3 mmol) in THF (20 mL) stirred at -78°C under nitrogen was dropwise added n-BuLi (2.5M in hexane, 1.89 mL, 4.72 mmol). The mixture was stirred for 2 hours at -78°C, then warmed up to 0°C and transferred via cannula over a solution of γ -butyrolactone (362 μ L, 4.72 mmol) in THF (10 mL) cooled at -78°C. The reaction mixture was stirred for 2 hours at -78°C, quenched by a slow addition of water (10 mL), and then extracted with ethyl acetate (3 x 50 mL). The combined organic layers were dried (MgSO₄) and concentrated. Purification by flash silica gel chromatography (40% ethyl acetate in hexane) afforded the corresponding the alcohol 200 (430 mg, 42% yield). MS (ESI) = 241 (MH⁺).

Step 2: 3-[(4-Biphenyl)carbonyl]propionaldehyde (201)

To a suspension of pyridinium chlorochromate (448 mg, 2.08 mmol), sodium acetate (170 mg, 2.08 mmol), Celite (200 mg) and molecular sieves 4Å (10 mg) in anhydrous dichloromethane (10 mL) at room temperature under nitrogen was slowly added a solution of alcohol 200 in dichloromethane (5 mL). The reaction mixture was stirred at room temperature for 30 minutes and then filtered through a short pad of Celite. The organic layer was dried (MgSO₄), concentrated, and the crude was purified by flash silica gel chromatography (40% ethyl acetate in hexane) affording the aldehyde 201 (160 mg, 73% yield). MS (ESI) = 239 (MH⁺).

Step 3: (E,E)-Ethyl-7-[(4-biphenyl)carbonyl]-2,4-heptadienoate (202)

To a solution of aldehyde 201 (180mg, 0.75 mmol) in THF (10 mL), stirred at 0°C under nitrogen, was added triethyl-4-phosphonocrotonate (220 μ L, 0.98 mmol) followed by sodium hydride (40 mg, 0.98 mmol). The mixture was allowed to warm up to room temperature in 2 hours and then quenched by adding water, and extracted with ethyl acetate. The combined organic layers were washed with brine, dried (MgSO₄) and concentrated. Purification by flash silica gel

chromatography (30% ethyl acetate in hexane) afforded the corresponding diene 202 (100 mg, 40% yield). MS (ESI) = 335 (MH⁺).

Step 4: (E,E)-7-[(4-Biphenyl)carbonyl]-2,4-heptadienoic acid (203)

5 Following the procedure described in Example 18, step 3, but substituting ester 202 for ester 49a, the title compound 203 was obtained in 80% yield. ¹H NMR: (300 MHz, CDCl₃:CD₃OD 5:1) δ: 7.89-7.95 (2 m, 2 H), 7.49-7.65 (m, 4 H), 7.28-7.40 (m, 3 H), 7.15 (dd, J = 15.3, 9.9 Hz, 1 H), 6.05-6.24 (m, 2 H), 5.69 (d, J = 15.6 Hz, 1 H), 3.1 (t, J = 7.2 Hz, 2 H), 2.50-2.60 (m, 2 H).

10 Step 5: (E,E)-N-(2-Aminophenyl)-7-[(4-biphenyl)carbonyl]-2,4-heptadienamide (204)

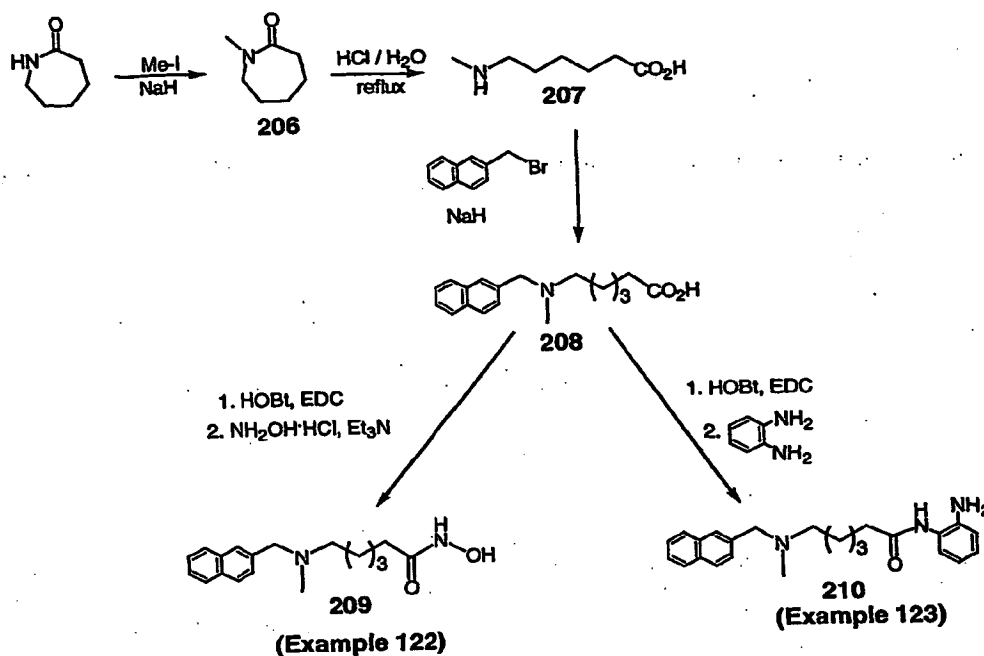
Following the procedure described in Example 22, but substituting carboxylic acid 203 for 50g, the title compound 204 was obtained in 47% yield. MS (ESI) = 397 (MH⁺).

Example 121:

15 (E,E)-N-Hydroxy-7-[(4-biphenyl)carbonyl]-2,4-heptadienamide (205)

Following the procedure described in Example 14, step 3, but substituting carboxylic acid 203 for 37, and using 1.1 equivalent of NH₂OH.HCl and triethylamine each, the title compound 205 was obtained in 35% yield. MS (ESI) = 322 (MH⁺).

20

**Example 122:****N-Hydroxy-N-(2-naphthylmethyl-methylamino)hexanamide (209)****Step 1: N-Methyl- ϵ -caprolactam (206)**

- 5 To a solution of ϵ -caprolactam (2.0 g, 17.6 mmol) in anhydrous THF (15 mL) at 0°C was added NaH, 60% dispersion in oil (1.06 g, 26.0 mmol), and the mixture was stirred at 0°C for 30 minutes. The cold bath was removed and the mixture was stirred at room temperature for 1 hour and then cooled again to 0°C.
- 10 Methyl iodide (1.65 mL, 26.0 mmol) was added dropwise over 10 minutes, then the cold bath was removed and the mixture was stirred at room temperature for 18 hours. The excess NaH was quenched with methanol, THF was removed in *vacuo* and the crude was partitioned between ethyl acetate and water. The aqueous phase was washed with ethyl acetate (2 x 30 mL), adding solid NaCl to break the emulsion. Organic washings were combined and dried over Na₂SO₄.
- 15 Ethyl acetate was removed in *vacuo* and the crude product 206 (1.54 g, 68%) was used without further purification. MS (ESI) = 128 (MH⁺).

Step 2: 6-N-Methylamino hexanoic acid (207)

A mixture of N-methyl ϵ -caprolactam 206 (1.1 g, 8.6 mmol) and 8N HCl (8 mL) and H₂O (4 mL) was refluxed for 18 hours. The resulting solution was cooled to room temperature, diluted with H₂O (20 mL) and the mixture was evaporated to dryness. Additional H₂O (10 mL) was added and again removed in *vacuo*. The crude residue was triturated with acetone (2 X 5 mL), removing acetone each time. The resulting solid was dried in *vacuo* to yield the desired product 207 (1.21 g, 96%). MS (ESI) = 146 (MH⁺).

Step 3: 6-N-(2-naphthylmethyl-methylamino)hexanoic acid (208)

Following the procedure described in Example 122, step 1, but substituting respectively the amino acid 207 for the azepan-2-one, 2-bromomethyl-naphthalene for methyl iodide, and DMF for THF, the title compound 208 was obtained in 45% yield after purification by column chromatography (33% to 90% methanol in dichloromethane).

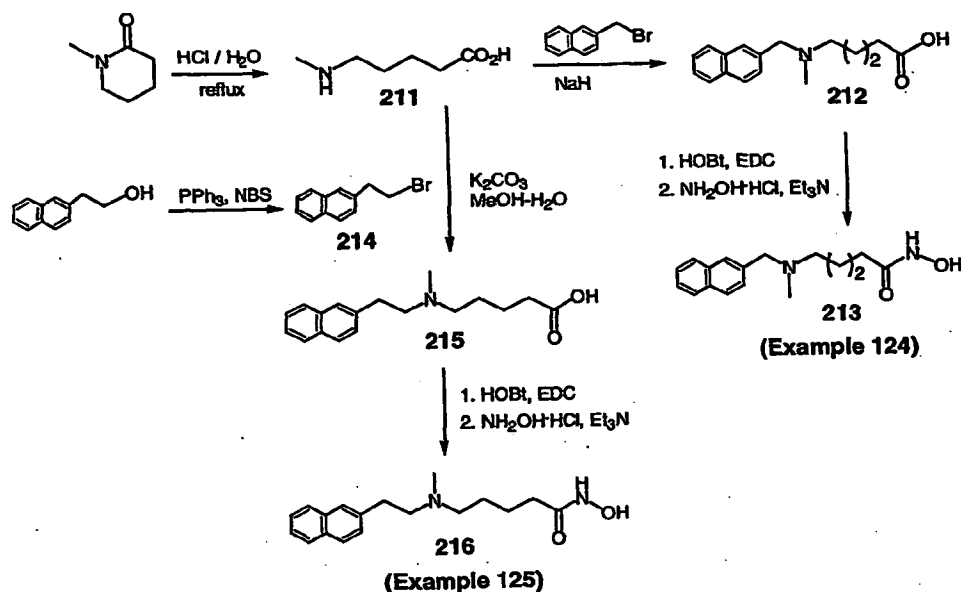
Step 4: N-Hydroxy-6-N-(2-naphthylmethyl-methylamino)hexanamide (209)

Following the procedure described in Example 14, step 3, but substituting carboxylic acid 208 for 37, and using 1.1 equivalent of NH₂OH · HCl and triethylamine each, the title compound 209 was obtained in 20% yield. ¹H NMR: (300 MHz, DMSO-d₆) δ 10.3 (br s, 1H), 8.65 (br d, J = 0.8 Hz, 1H), 7.86 (m, 3H), 7.80 (s, 1H), 7.47 (m, 3H), 3.58 (s, 2H), 2.33 (t, J = 7.2 Hz, 2H), 2.12 (s, 3H), 1.93 (t, J = 7.3 Hz, 2H), 1.48 (m, 4H), 1.26 (m, 2H).

Example 123:

N-(2-Aminophenyl)-N-(2-naphthylmethyl-methylamino)hexanamide (210)

Following the procedure described in Example 22, but substituting carboxylic acid 208 for 50g, the title compound 210 was obtained in 20% yield. ¹H NMR: (300 MHz, DMSO-d₆) δ 9.1 (br s, 1H), 7.86 (m, 3H), 7.86 (m, 3H), 7.78 (br s, 1H), 7.47 (m, 3H), 7.14 (dd, J = 8.0, 1.4 Hz, 1H), 6.87 (dt, J = 8.0, 1.4 Hz, 1H), 6.70 (dd, J = 8.0, 1.4 Hz, 1H), 6.51 (dt, J = 7.4, 1.4 Hz, 1H), 4.81 (br, 2H), 3.63 (br, 2H), 2.39 (br, 2H), 2.30 (t, J = 7.3 Hz, 2H), 2.16 (br s, 3H), 1.56 (m, 4H), 1.35 (m, 2H).

**Example 124:****N-Hydroxy-5-N-(2-naphthylmethyl-methylamino)pentanamide (213)****Step 1: 5-Methylamino-pentanoic acid (211)**

5 Following the procedure described in Example 122, step 2, but substituting 1-methyl-2-piperidone for 206, the title compound 211 was obtained in 95% yield. MS (ESI) = 132 (MH^+).

Step 2: 5-N-(2-naphthylmethyl-methylamino)pentanoic acid (212)

10 Following the procedure described in Example 122, step 3, but substituting amino acid 211 for 207, the title compound 212 was obtained in 58% yield. MS (ESI) = 272 (MH^+).

Step 3: N-Hydroxy-5-N-(2-naphthylmethyl-methylamino)pentanamide (213)

15 Following the procedure described in Example 14, step 3, but substituting carboxylic acid 212 for 37, and using 1.1 equivalent of $\text{NH}_2\text{OH} \cdot \text{HCl}$ and triethylamine each, the title compound 213 was obtained in 33% yield. ^1H NMR: (300 MHz, $\text{DMSO}-d_6$) δ 10.30 (br s, 1H), 8.61 (br s, 1H), 7.85 (m, 3H), 7.75 (s, 1H),

7.46 (m, 3H), 3.57 (s, 2H), 2.34 (t, J = 6.9 Hz, 2H), 2.1 (s, 3H), 1.93 (t, J = 6.9 Hz, 2H), 1.47 (m, 4H).

Example 125:

N-Hydroxy-5-N-[2-(2-naphthylethyl)-methylaminol]pentanamide (216)

5 **Step 1: 2-(2-Bromoethyl)naphthalene (214)**

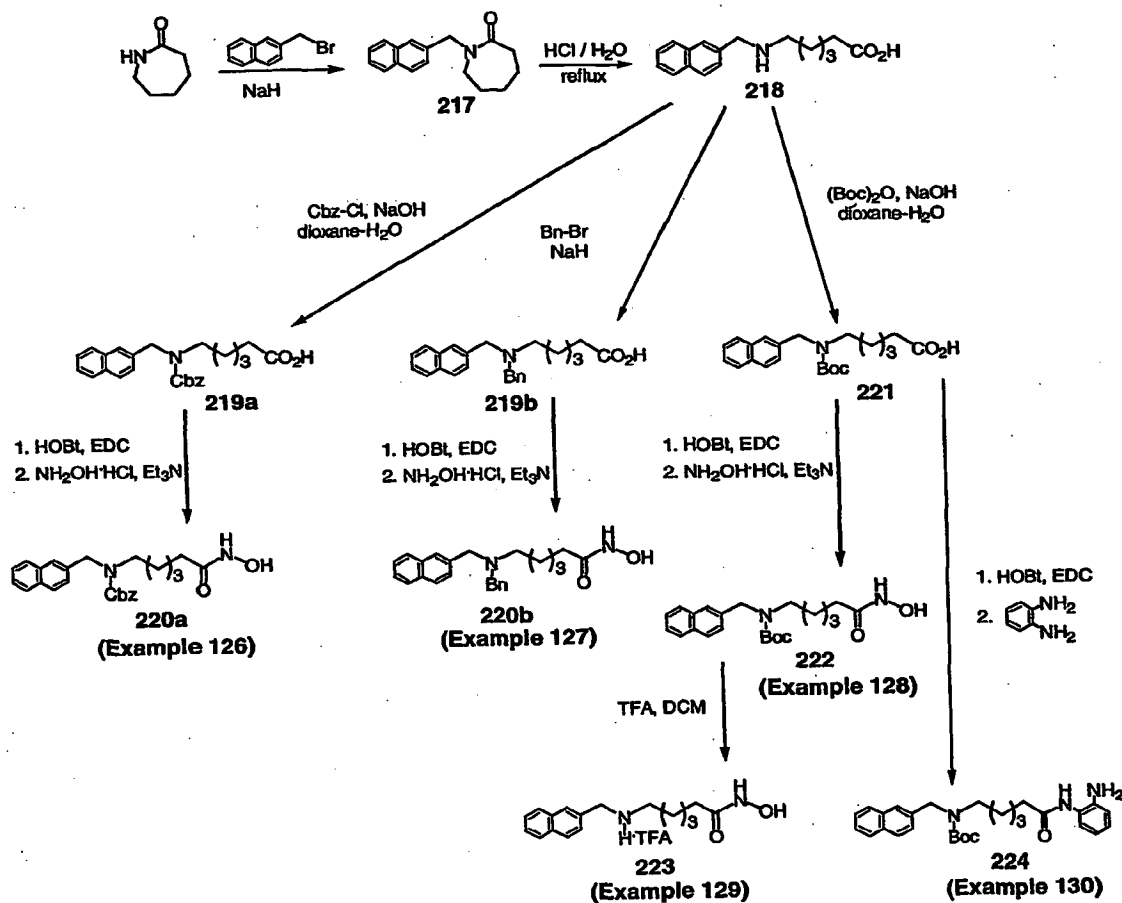
A solution of triphenylphosphine (0.57 g, 2.2 mmol) and N-bromosuccinimide (0.39 g, 2.2 mmol) in anhydrous dichloromethane (4 mL) was stirred for 10 minutes. 2-Naphthalenethanol (0.25 g, 1.45 mmol) was added, followed immediately by imidazole (0.25 g, 1.45 mmol) and the mixture was stirred at room temperature for 18 hours. The reaction mixture was partitioned between H₂O and dichloromethane. The aqueous layer was extracted with diethyl ether (2 x 20 mL) and the combined organic extracts were dried over MgSO₄ and evaporated to dryness. The crude residue was purified by column chromatography through a plug of silica gel eluting with 10% ethyl acetate in hexanes to afford a colorless solid 214 (0.29 g, 86%).

Step 2: 5-N-[2-(2-Naphthylethyl)-methylaminol]pentanoic acid (215)

A mixture of amino acid 211 (60 mg, 0.45 mmol), 2-(2-bromoethyl)naphthalene (0.15 g, 0.64 mmol) and potassium carbonate (0.24 g, 1.7 mmol) in methanol-H₂O (2 : 1.5 mL) was heated at reflux for 7 hours and then at room temperature for 18 hours. Methanol was removed in *vacuo*, the remaining aqueous residue was partitioned between ethyl acetate and H₂O and extracted with ethyl acetate (2 x 20 mL). The aqueous layer was evaporated to dryness, triturated with methanol and filtered to remove inorganic solids. The filtrate was evaporated to dryness and then purified by column chromatography using a gradient of 25 to 75% methanol in dichloromethane to afford a colorless solid 215 (85 mg, 69%). MS (ESI) = 286 (M+1).

Step 3: *N*-Hydroxy-5-*N*-[2-(2-naphthylethyl)-methylamino]pentanamide (216)

Following the procedure described in Example 14, step 3, but substituting carboxylic acid 215 for 37, and using 1.1 equivalent of $\text{NH}_2\text{OH}\cdot\text{HCl}$ and triethylamine each, the title compound 216 was obtained in 15% yield. ^1H NMR: (300 MHz, $\text{DMSO}-d_6$) δ 7.83 (m, 3H), 7.71 (s, 1H), 7.44 (m, 3H), 2.86 (dd, $J = 8.5, 6.9$ Hz, 2H), 2.60 (m, 2H), 2.34 (t, $J = 7.0$ Hz, 2H), 2.2 (s, 3H), 1.93 (t, $J = 7.15$ Hz, 2H), 1.43 (m, 4H).



Example 126:**N-Hydroxy-6-N-(Benzyloxycarbonyl-2-naphthylmethylamino)hexanamide (220a)****Step 1: N-2-Naphthylmethyl-ε-caprolactam (217)**

5 Following the procedure described in Example 122, step 1, but substituting respectively 2-bromomethylnaphthalene for methyl iodide, and DMF for THF, the title compound 217 was obtained in 97% yield. MS (ESI) = 254 (MH⁺).

Step 2: 6-N-(2-Naphthylmethylamino)hexanoic acid (218)

10 Following the procedure described in Example 122, step 2, but substituting 217 for 206, the title compound 218 was obtained in 69% yield. MS (ESI) = 272 (MH⁺).

Step 3: 6-N-(Benzyloxycarbonyl-2-naphthylmethylamino)hexanoic acid (219a)

15 To an emulsion of compound 218 (70 mg, 0.26 mmol) in dioxane-H₂O (1.5 : 2.5 mL) stirring at room temperature was added NaOH (0.1 g, 2.5 mmol) dissolved in 1 mL H₂O followed by benzyloxycarbonyl chloride (0.044 mL, 0.31 mmol). The mixture was stirred vigorously for 18 hours. Dioxane was removed in *vacuo* and the resulting aqueous layer was diluted with H₂O and extracted with diethyl ether (2 x 20mL). Subsequently, the aqueous layer was neutralized to pH 7 using a KHSO₄ solution (0.37 M) and then extracted with ethyl acetate (2 x 30 mL).
20 The combined organic extracts were dried over Na₂SO₄ and then evaporated to dryness to afford a crude oil 219a (75 mg, 72%) that was used without further purification. MS (ESI) = 428 (MH⁺).

Step 4: N-Hydroxy-6-N-(Benzyloxycarbonyl-2-naphthylmethylamino)hexanamide (220a)

25 Following the procedure described in Example 14, step 3, but substituting carboxylic acid 219a for 37, and using 1.1 equivalent of NH₂OH.HCl and triethylamine each, the title compound 220a was obtained in 66% yield. ¹H NMR: (300 MHz, DMSO-d₆) δ 10.30 (br s, 1H), 8.65 (br s, 1H), 7.87 (m, 3H), 7.7 (br d, J = 10.7 Hz, 1H), 7.49 (t, J = 3.7 Hz, 2H), 7.38 (m, 4H), 7.25 (m, 2H), 5.13 (br d, J = 6.6

Hz, 2H), 4.61 (s, 2H), 3.22 (m, 2H), 2.1 (m, 1H), 1.88 (m, 2H), 1.47 (m, 4H), 1.20 (m, 2H).

Example 127:

***N*-Hydroxy-6-*N*-(Benzyl-2-naphthylmethylamino)hexanamide (220b)**

5 **Step 1: 6-*N*-(Benzyl-2-naphthylmethylamino)hexanoic acid (219b)**

Following the procedure described in Example 122, step 1, but substituting respectively the amino acid 218 for the ϵ -caprolactam, benzyl bromide for methyl iodide, and DMF for THF, the title compound 219b was obtained in 56% yield. MS (ESI) = 362 (MH⁺).

10 **Step 2: *N*-Hydroxy-6-*N*-(benzyl-2-naphthylmethylamino)hexanamide (220b)**

Following the procedure described in Example 14, step 3, but substituting carboxylic acid 219b for 37, and using 1.1 equivalent of NH₂OH.HCl and triethylamine each, the title compound 220b was obtained in 27% yield. ¹H NMR: (300 MHz, DMSO-d₆) δ 10.29 (br s, 1H), 8.65 (br s, 1H), 7.84 (m, 4H), 7.48 (m, 3H),
15 7.33 (m, 4H), 7.21 (m, 1H), 3.66 (s, 2H), 3.54 (s, 2H), 3.35 (t, J = 7.1 Hz, 2H), 1.87 (t, J = 7.3 Hz, 2H), 1.46 (m, 2H), 1.40 (m, 2H), 1.19 (m, 2H).

Example 128:

***N*-Hydroxy-6-*N*-(*tert*-butoxycarbonyl-2-naphthylmethylamino)hexanamide (222)**

20 **Step 1: 6-*N*-(*tert*-Butoxycarbonyl-2-naphthylmethylamino)hexanoic acid (221)**

Following the procedure described in Example 126, step 3, but substituting di-*tert*-butyl dicarbonate for benzyloxycarbonyl chloride, the title compound 221 was obtained in 88% yield. MS (ESI) = 394 (MH⁺).

25 **Step 2: *N*-Hydroxy-6-*N*-(*tert*-butoxycarbonyl-2-naphthylmethylamino)hexanamide (222)**

Following the procedure described in Example 14, step 3, but substituting carboxylic acid 221 for 37, and using 1.1 equivalent of NH₂OH.HCl and

triethylamine each, the title compound **222** was obtained in 60% yield. ^1H NMR : (300 MHz, DMSO- d_6) δ 10.31 (br s, 1H), 8.65 (br s, 1H), 7.87 (m, 3H), 7.7 (br s, 1H), 7.5 (m, 2H), 7.37 (m, 1H), 4.52 (br s, 2H), 3.15 (m, 2H), 1.90 (t, $J = 7.3$ Hz, 2H), 1.41 (m, 13H), 1.17 (m, 2H).

5 **Example 129:**

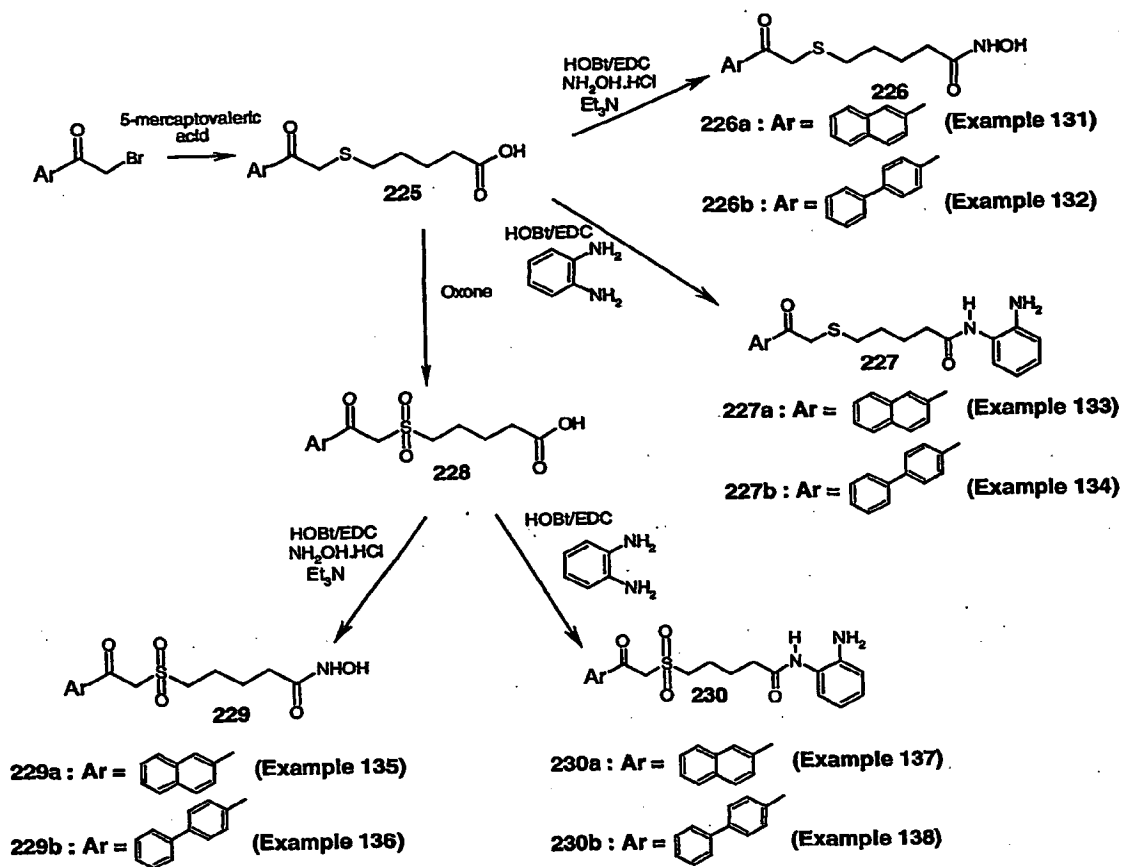
***N*-Hydroxy-6-*N*-(2-naphthylmethylamino)hexanamide trifluoroacetate (**223**)**

To a stirring solution of compound **222** (65.0 mg, 0.17 mmol) in anhydrous dichloromethane (3 mL) at room temperature was added trifluoroacetic acid (3.5 mL) dropwise. The solution was stirred for 18 hours. dichloromethane and
10 trifluoroacetic acid were removed in *vacuo*, the residue was dissolved in H_2O and then extracted with ethyl acetate (2 x 20 mL). The aqueous layer was lyophilized to yield the final product **223** as a colorless oil (30.0 mg, 45%). ^1H NMR : (300 MHz, DMSO- d_6) δ 10.35 (s, 1H), 8.79 (br s, 2H), 8.67 (s, 1H), 7.96 (m, 4H), 7.58 (m, 3H), 4.3 (br s, 2H), 2.94 (br s, 2H), 1.93 (t, $J = 7.4$ Hz, 2H), 1.61 (m, 2H), 1.49 (m,
15 2H), 1.27 (m, 2H).

Example 130:

***N*-(2-Aminophenyl)-6-*N*-(*tert*-butoxycarbonyl-2-naphthylmethylamino)-hexanamide (**224**)**

Following the procedure described in Example 22, but substituting
20 carboxylic acid **221** for 50g, the title compound **224** was obtained in 53% yield. ^1H NMR: (300 MHz, DMSO- d_6) δ 9.06 (br s, 1H), 7.87 (m, 3H), 7.71 (br s, 1H), 7.49 (m, 2H), 7.38 (d, $J = 8.5$ Hz, 1H), 7.13 (d, $J = 6.6$ Hz, 1H), 6.88 (m, 1H), 6.70 (dd, $J = 1.1$, 8.0 Hz, 1H), 6.52 (dt, $J = 1.4$, 8.2 Hz, 1H), 4.8 (m, 2H), 4.54 (br s, 2H), 3.17 (m, 2H), 2.28 (t, $J = 7.3$ Hz, 2H), 1.44 (m, 15H).

**Example 131:*****N*-Hydroxy-5-(2-naphthoylethylsulfanyl)pentanamide (226a)****Step 1: 5-(2-Naphthoylethylsulfanyl)pentanoic acid (225a)**

- 5 To a solution of bromomethyl-2-naphthyl ketone (2 g, 8.00 mmol) in anhydrous THF (50 mL) was added 5-mercaptovaleric acid (1.07 g, 8.00 mmol), followed by potassium carbonate (5.71 g, 41 mmol). The suspension was refluxed for 1 hour, cooled to the room temperature and filtered. The solid was collected, dissolved in a 1:1 mixture water-THF and acidified with HCl (pH 1-2). The acidic
- 10 solution was extracted with ether, dried (MgSO₄), filtered and evaporated to produce the title compound 225a (1.21 g, 50% yield). ¹H NMR: (300 MHz, DMSO-d₆) δ 11.98 (br s, 1H), 8.69 (s, 1H), 8.10 (d, J = 7.5 Hz, 1H), 8.00-7.98 (m, 3H), 7.67-7.61 (m, 2H), 4.09 (s, 2H), 2.52 (t, J = 6.6 Hz, 2H), 2.19 (t, J = 6.6 Hz, 2H), 1.56-1.53 (m, 4H).

Step 2: N-Hydroxy 5-(2-naphthoylmethylsulfanyl)pentanamide (226a)

Following the procedure described in Example 14, step 3, but substituting carboxylic acid 225a for 37, and using 1.1 equivalent of $\text{NH}_2\text{OH}\cdot\text{HCl}$ and triethylamine each, the title compound 226a was obtained in 24% yield. ^1H NMR: (300 MHz, DMSO-d_6) δ 10.33 (s, 1H), 8.70 (s, 1H), 8.66 (s, 1H), 8.10 (d, $J = 7.7$ Hz, 1H), 8.00-7.97 (m, 3H), 7.70-7.59 (m, 2H), 2.51 (t, $J = 6.9$ Hz, 2H), 4.10 (s, 2H), 1.94-1.91 (m, 2H), 1.53-1.51 (m, 4H).

Example 132:

N-Hydroxy-5-(4-biphenylcarbonylmethanesulfanyl)pentanamide (226b)

Step 1: 5-(4-Biphenylcarbonylmethanesulfanyl)pentanoic acid (225b)

Following the procedure described in Example 131, step 1, but substituting bromomethyl-4-biphenyl ketone for bromomethyl-2-naphthyl ketone, the title compound 225b was obtained in 81% yield. ^1H NMR: (300 MHz, CDCl_3) δ 8.05 (d, $J = 8.0$ Hz, 2H), 7.71-7.62 (m, 4H), 7.50-7.40 (m, 3H), 3.81 (s, 2H), 2.60 (t, $J = 6.3$ Hz, 2H), 2.38 (t, $J = 6.9$ Hz, 2H), 1.72-1.68 (m, 4H).

Step 2: N-Hydroxy-5-(4-biphenylcarbonylmethanesulfanyl)pentanamide (226b)

Following the procedure described in Example 14, step 3, but substituting carboxylic acid 225b for 37, and using 1.1 equivalent of $\text{NH}_2\text{OH}\cdot\text{HCl}$ and triethylamine each, the title compound 226b was obtained in 30% yield. ^1H NMR: (300 MHz, DMSO-d_6) δ 10.31 (s, 1H), 8.65 (s, 1H), 8.05 (d, $J = 8.2$ Hz, 2H), 7.73 (d, $J = 8.0$ Hz, 2H), 7.80 (d, $J = 8.5$ Hz, 2H), 7.51-7.41 (m, 3H), 3.98 (s, 2H), 1.92 (br s, 4H), 1.51 (br. s, 4H),

Example 133:

N-(2-Aminophenyl)-5-(2-naphthoylmethylsulfanyl)pentanamide (227a)

Following the procedure described in Example 22, but substituting carboxylic acid 225a for 50g, the title compound 227a was obtained in 20% yield. ^1H NMR: (300 MHz, DMSO-d_6) δ 9.07 (s, 1H), 8.69 (s, 1H), 8.10-7.97 (m, 4H), 7.68-

7.58 (m, 2H), 7.11 (d, $J = 7.1$ Hz, 1H), 6.86 (t, $J = 7.1$ Hz, 1H), 6.80 (d, $J = 7.7$ Hz, 1H), 6.50 (t, $J = 7.4$ Hz, 1H), 4.78 (s, 2H), 4.10 (s, 2H), 2.56 (t, $J = 6.6$ Hz, 2H), 2.30 (t, $J = 6.6$ Hz, 2H), 1.61 (br s, 4H).

Example 134:

5 ***N*-(2-Aminophenyl)-5-(4-biphenylcarbonylmethanesulfanyl)pentanamide (227b)**

Following the procedure described in Example 22, but substituting carboxylic acid 225b for 50g, the title compound 227b was obtained in 34% yield.

¹H NMR: (300 MHz, DMSO-*d*₆) δ 9.07 (s, 1H), 8.05 (d, $J = 8.2$ Hz, 2H), 7.80 (d, $J = 8.0$ Hz, 2H), 7.73 (d, $J = 7.7$ Hz, 2H), 7.51-7.41 (m, 3H), 7.12 (d, $J = 7.5$ Hz, 1H), 6.86 (t, $J = 7.5$ Hz, 1H), 6.69 (d, $J = 7.8$ Hz, 1H), 6.51 (t, $J = 7.5$ Hz, 1H), 4.79 (s, 2H), 4.00 (s, 2H), 2.53 (t, $J = 6.6$ Hz, 2H), 2.30 (t, $J = 7.2$ Hz, 2H), 1.61 (br s, 4H).

Example 135:

***N*-Hydroxy-5-(2-naphthoylmethylsulfonyl)pentanamide (229a)**

15 **Step 1: 5-(2-Naphthoylmethylsulfonyl)pentanoic acid (228a)**

To a solution of 225a (755 mg, 2.50 mmol) in methanol (30 mL) at 0°C was added a solution of oxone (2.3 g, 3.75 mmol) in water (30 mL). The mixture stirred at the same conditions for 6 hours, methanol was removed in vacuum and the water phase was extracted with chloroform. The extract was dried over MgSO₄,
20 filtered and evaporated to afford the title compound 228a (520 mg, 62% yield). MS (ESI) = 335 (MH⁺).

Step 2: *N*-Hydroxy-5-(2-naphthoylmethylsulfonyl)pentanamide (229a)

Following the procedure described in Example 14, step 3, but substituting
25 carboxylic acid 228a for 37, and using 1.1 equivalent of NH₂OH.HCl and triethylamine each, the title compound 229a was obtained in 30% yield. ¹H NMR: (300 MHz, DMSO-*d*₆) δ 10.38 (br s, 1H), 8.83 (s, 1H), 8.71 (s, 1H), 8.13 (d, $J = 7.7$ Hz,

1H), 8.06-7.99 (m, 3H), 7.74-7.63 (m, 2H), 5.19 (s, 1H), 3.34 (t, J = 7.4 Hz, 2H), 2.00 (t, J = 6.7 Hz, 2H), 1.78-1.60 (m, 4H).

Example 136:

N-Hydroxy-5-(4-biphenylcarbonylmethylsulfonyl)pentanamide (229b)

5 **Step 1: 5-(4-Biphenylcarbonylmethanesulfonyl)pentanoic acid (228b)**

Following the procedure described in Example 135, step 1, but substituting 228b for 228a, the title compound 228b was obtained in 91% yield.

Step 2: N-Hydroxy-5-(4-biphenylcarbonylmethylsulfonyl)pentanamide (229b)

10 Following the procedure described in Example 14, step 3, but substituting carboxylic acid 228b for 37, and using 1.1 equivalent of $\text{NH}_2\text{OH}\cdot\text{HCl}$ and triethylamine each, the title compound 228b was obtained in 52% yield. ^1H NMR: (300 MHz, DMSO-d_6) δ 10.37 (s, 1H), 8.70 (s, 1H), 8.11 (d, J = 8.5 Hz, 2H), 7.86 (d, J = 8.5 Hz, 2H), 7.77 (dd, J = 1.4 Hz, J = 8.5 Hz, 2H), 7.53-7.43 (m, 3H), 5.08 (s, 2H), 3.38-3.25 (t, 2H), 1.99 (t, J = 6.6 Hz, 2H), 1.72-1.61 (m, 4H).

15 **Example 137:**

N-(2-Aminophenyl)-5-(2-naphthoylmethylsulfonyl)pentanamide (230a)

Following the procedure described in Example 22, but substituting carboxylic acid 228a for 50g, the title compound 230a was obtained in 30% yield. ^1H NMR: (300 MHz, DMSO-d_6) δ 9.12 (s, 1H), 8.83 (s, 1H), 8.12 (d, J = 7.4 Hz, 1H), 8.06-7.99 (m, 3H), 7.74-7.63 (m, 2H), 7.15 (dd, J = 1.4 Hz, J = 8.0 Hz, 1H), 6.88 (dd, J = 1.4 Hz, 8.0 Hz, 1H), 6.70 (dd, J = 1.4 Hz, 8.0 Hz, 1H), 6.52 (dd, J = 1.2 Hz, 8.8 Hz, 1H), 5.21 (s, 1H), 4.82 (br s, 2H), 3.38 (t, J = 7.1 Hz, 2H), 2.38 (t, J = 7.1 Hz, 2H), 1.84-1.72 (m, 4H).

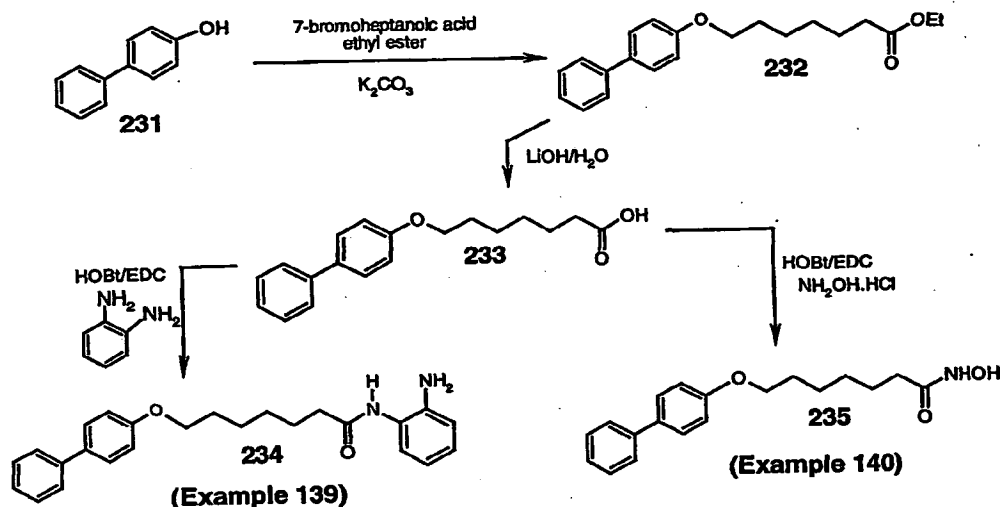
Example 138:

25 **N-(2-Aminophenyl)-5-(4-biphenylcarbonylmethylsulfonyl)pentanamide (229b)**

Following the procedure described in Example 22, but substituting carboxylic acid 228b for 50g, the title compound 230b was obtained in 52% yield.

¹H NMR: (300 MHz, DMSO-d₆) δ 9.12 (s, 1H), 8.13 (d, J = 8.2 Hz, 2H), 7.87 (d, J = 8.2 Hz, 2H), 7.77 (d, J = 6.7 Hz, 2H), 7.53-7.44 (m, 3H), 7.15 (d, J = 8.0 Hz, 1H), 6.88 (t, J = 8.0 Hz, 1H), 6.70 (d, J = 8.0 Hz, 1H), 6.52 (t, J = 7.7 Hz, 1H), 5.11 (s, 2H), 4.82 (s, 2H), 3.36 (t, J = 6.9 Hz, 2H), 2.37 (t, J = 7.1 Hz, 2H), 1.78-1.71 (m, 4H).

5



Example 139:

N-(2-Aminophenyl)-7-(4-biphenyloxy)heptanamide (234)

Step 1: Ethyl-7-(4-biphenyloxy)heptanoate (232)

10 To a solution of 4-phenylphenol (231) (1.00 g, 5.88 mmol) in acetone (100 mL) was added potassium carbonate (3.25 g, 23.52 mmol) and potassium iodide (200 mg, 1.57 mmol), followed by the addition of ethyl-7-bromoheptanoate (1.81 g, 7.65 mmol). The suspension was refluxed for 36 hours, cooled to room temperature and filtered. Filtrate was evaporated and the remained solid was
 15 recrystallized from hexane to produce the title compound 232 (1.73 g, 90% yield). MS (ESI) = 327 (MH⁺).

Step 2: 7-(4-Biphenyloxy)heptanoic acid (233)

To a solution of the ester 232 (1.70 g, 5.21 mmol) in THF (50 mL) was added a solution of lithium hydroxide hydrate (1.30 g, 30.95 mmol) in water (50 mL). The

reaction mixture was stirred 18 hours at room temperature, THF was evaporated and the aqueous phase was acidified with conc. HCl (pH 1-2). A precipitate formed which was collected by filtration and dried to afford the title compound 233 (1.54 g, 99%). MS (ESI) = 321 ($M^+ + Na$).

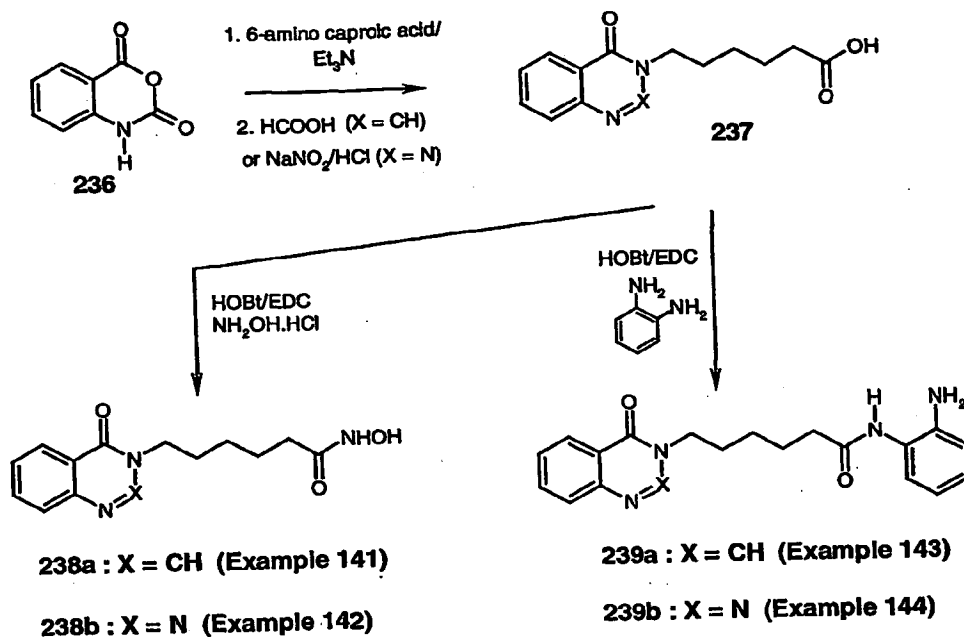
5 Step 3: N-(2-Aminophenyl)-7-(4-biphenyloxy)heptanamide (234)

Following the procedure described in Example 22, but substituting carboxylic acid 233 for 50g, the title compound 234 was obtained in 44% yield. 1H NMR: (300 MHz, DMSO- d_6) δ 9.08 (s, 1H), 7.60-7.55 (m, 4H), 7.41 (t, J = 7.7 Hz, 2H), 7.29 (t, J = 7.1 Hz, 1H), 7.14 (dd, J = 8.0, 1.4 Hz, 1H), 7.00 (d, J = 8.8 Hz, 2H),
10 6.87 (t, J = 8.0 Hz, 1H) 6.70 (dd, J = 8.0, 1.4 Hz, 1H), 6.52 (dd, J = 8.5, 1.1 Hz, 1H), 4.81 (br s, 2H), 4.00 (t, J = 6.3 Hz, 2H), 2.32 (t, J = 7.4 Hz, 2H), 1.76-1.69 (m, 2H), 1.67-1.57 (m, 2H), 1.47-1.38 (m, 4H).

Example 140:

N-Hydroxy-7-(4-biphenyloxy)heptanamide (235)

15 Following the procedure described in Example 14, step 3, but substituting carboxylic acid 233 for 37, and using 1.1 equivalent of $NH_2OH \cdot HCl$ and triethylamine each, the title compound 235 was obtained in 52% yield. 1H NMR: (300 MHz, DMSO- d_6) δ 10.33 (s, 1H), 8.65 (s, 1H), 7.60-7.54 (m, 4H), 7.40 (t, J = 7.5 Hz, 2H), 7.28 (t, J = 7.2 Hz, 1H), 6.98 (d, J = 8.7 Hz, 2H), 3.97 (t, J = 6.3 Hz, 2H), 1.93
20 (t, J = 7.1 Hz, 2H), 1.70 (t, J = 7.4 Hz, 2H), 1.52-1.28 (m, 6H).

**Example 141:*****N*-Hydroxy-6-[3-(4-Oxo-4*H*-quinazolinyl)]hexanamide (238a)****Step 1: 6-[3-(4-Oxo-4*H*-quinazolinyl)]hexanoic acid (237a)**

5 To a solution of 6-aminohexanoic acid (2.88 g, 20.00 mmol) in water (50 mL) was added triethylamine (3.06 mL, 22.00 mmol) followed by a portionwise addition of 1*H*-benzo[d][1,3]oxazine-2,4-dione (isatoic anhydride, 236) (3.26 g, 20.00 mmol). The reaction mixture was stirred for 2 hours at 30–40°C, cooled to room temperature and evaporated in vacuum to form an oily residue. This

10 material was refluxed for 7 hours in formic acid (70 mL), cooled to room temperature and evaporated. The solid was dissolved in dichloromethane, decolorized with the activated charcoal, filtered and evaporated to give an oily material which upon crystallization in methanol (–10°C) afforded the title compound 237a (2.38 g, 46% yield). MS (ESI) = 261 (MH⁺).

Step 2: *N*-Hydroxy-6-[3-(4-oxo-4*H*-quinazolinyl)]hexanamide (238a)

15 Following the procedure described in Example 14, step 3, but substituting carboxylic acid 1*H*-benzo[d][1,3]oxazine-2,4-dione 237a for 37, and using 1.1 equivalent of NH₂OH.HCl and triethylamine each, the title compound 238a was

obtained in 23% yield. ¹H NMR: (400 MHz, DMSO-d₆) δ 10.32 (s, 1H), 8.66 (s, 1H), 8.38 (s, 1H), 8.14 (br d, J = 7.1 Hz, 1H), 7.81 (dd, J = 8.2, 1.3 Hz, 1H), 7.66 (br d, J = 8.1 Hz, 1H), 7.53 (br t, J = 7.7 Hz, 1H), 3.95 (t, J = 7.1 Hz, 2H), 1.93 (t, J = 7.2 Hz, 2H), 1.72-1.62 (m, 2H), 1.56-1.46 (m, 2H), 1.30-1.23 (m, 2H).

5 **Example 142:**

N-Hydroxy-6-[3-(4-oxo-4H-benzo[d][1,2,3]triazinyl)]hexanamide (238b)

Step 1: 6-[3-(4-Oxo-4H-benzo[d][1,2,3]triazinyl)]hexanoic acid (237b)

To a solution of 6-aminohexanoic acid (2.88 g, 20.00 mmol) in water (50 mL) was added triethylamine (3.06 mL, 22.00 mmol) followed by a portionwise
10 addition of 1H-benzo[d][1,3]oxazine-2,4-dione (isatoic anhydride, 236) (3.26 g, 20.00 mmol). The reaction mixture was stirred for 2 hours at 30-40°C, cooled to 0°C, acidified with 20 mL 20% HCl and treated with a solution of sodium nitrite (1.70 g, 25 mmol) in 10 mL water (over a 10 min period of time). The whole
15 mixture was stirring at ambient temperature for 18 hours to form a solid which was collected by filtration, dried, dissolved in dichloromethane and decolorized with the activated charcoal, filtered and evaporated. The residual material was crystallized in a dichloromethane-hexane mixture at -10°C to afford the title compound 237b in 75% yield. MS (ESI) = 262 (MH⁺).

Step 2: N-Hydroxy-6-[3-(4-oxo-4H-benzo[d][1,2,3]triazinyl)]hexanamide (238b)

20 Following the procedure described in Example 14, step 3, but substituting carboxylic acid 237b for 37, and using 1.1 equivalent of NH₂OH.HCl and triethylamine each, the title compound 238b was obtained in 32%. ¹H NMR: (400 MHz, DMSO-d₆) δ 10.32 (s, 1H), 8.66 (s, 1H), 8.25 (dd, J = 7.9, 1.0 Hz, 1H), 8.19 (br
25 d, J = 7.7 Hz, 1H), 8.08 (dd, J = 7.2, 1.4 Hz, 1H), 7.94 (dd, J = 7.2, 1.4 Hz, 1H), 4.37 (t, J = 7.1 Hz, 2H), 1.94 (t, J = 7.2 Hz, 2H), 1.85-1.75 (m, 2H), 1.58-1.48 (m, 2H), 1.35-1.27 (m, 2H).

Example 143:***N*-(2-Aminophenyl)-6-[3-(4-oxo-4*H*-quinazolinyl)]hexanamide (239a)**

Following the procedure described in Example 22, but substituting carboxylic acid 237a for 50g, the title compound 239a was obtained in 25% yield.

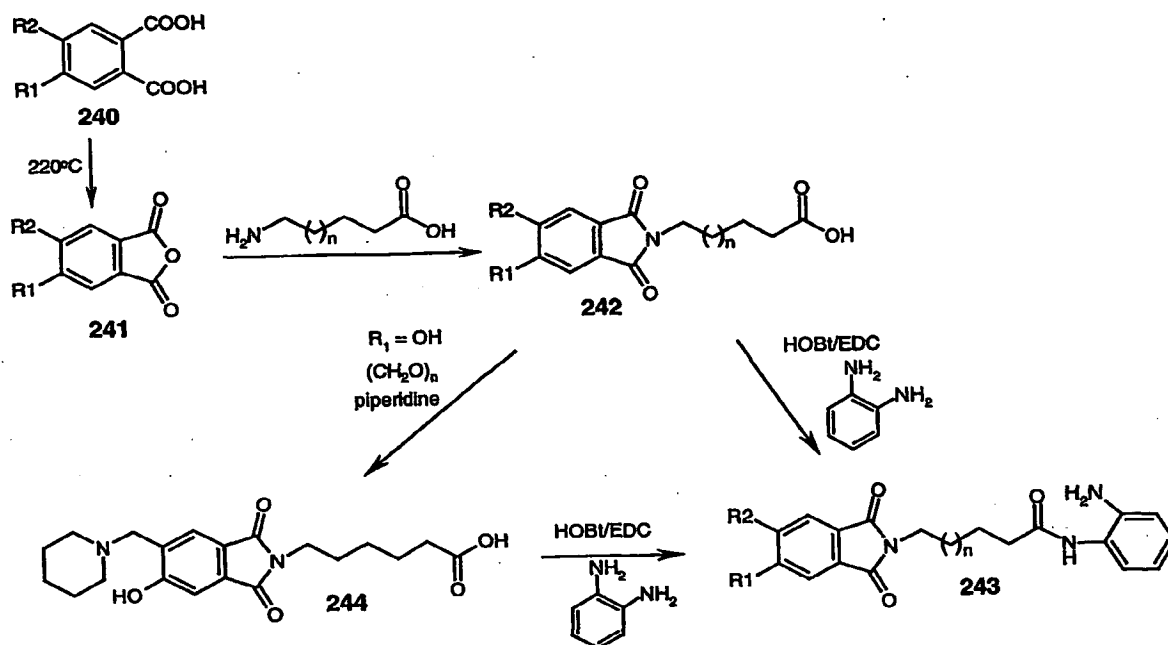
- 5 ¹H NMR: (400 MHz, DMSO-*d*₆) δ 9.08 (s, 1H), 8.41 (s, 1H), 8.17 (dd, *J* = 8.0, 1.1 Hz, 1H), 7.83 (dd, *J* = 8.4, 1.5 Hz, 1H), 7.68 (br d, *J* = 8.1 Hz, 1H), 7.55 (dd, *J* = 8.0, 1.0 Hz, 1H), 7.12 (dd, *J* = 7.8, 1.1 Hz, 1H), 6.88 (dd, *J* = 8.0, 1.3 Hz, 1H), 6.70 (dd, *J* = 7.9, 1.1 Hz, 1H), 6.51 (dd, *J* = 7.7, 1.3 Hz, 1H), 4.80 (s, 2H), 3.99 (t, *J* = 7.3 Hz, 2H), 2.32 (t, *J* = 7.3 Hz, 2H), 1.77-1.70 (m, 2H), 1.67-1.60 (m, 2H), 1.39-1.33 (m, 2H).

10 **Example 144:**

***N*-(2-Aminophenyl)-6-[3-(4-oxo-4*H*-benzo[*d*][1,2,3]triazinyl)]hexanamide (239b)**

Following the procedure described in Example 22, but substituting carboxylic acid 237b for 50g, the title compound 239b was obtained in 45% yield.

- 15 ¹H NMR: (400 MHz, DMSO-*d*₆) δ 9.07 (s, 1H), 8.25 (dd, *J* = 7.9, 0.9 Hz, 1H), 8.19 (br d, *J* = 8.1 Hz, 1H), 8.08 (dd, *J* = 7.2, 1.4 Hz, 1H), 7.92 (dd, *J* = 8.2, 1.2 Hz, 1H), 7.09 (dd, *J* = 7.8, 1.2 Hz, 1H), 6.87 (dd, *J* = 8.5, 1.4 Hz, 1H), 6.69 (dd, *J* = 8.0, 1.3 Hz, 1H), 6.49 (dd, *J* = 7.8, 1.4 Hz, 1H), 4.78 (s, 2H), 4.39 (t, *J* = 7.1 Hz, 2H), 2.28 (t, *J* = 7.4 Hz, 2H), 1.90-1.82 (m, 2H), 1.69-1.58 (m, 2H), 1.43-1.34 (m, 2H).



243a : $n = 2$, $\text{R}_1 = \text{R}_2 = \text{H}$ (Example 145)

243b : $n = 1$, $\text{R}_1 = \text{R}_2 = \text{H}$ (Example 146)


243c : $n = 3$, $\text{R}_1 = \text{R}_2 = \text{H}$ (Example 147)

243d : $n = 2$, $\text{R}_1 = \text{CH}_3$, $\text{R}_2 = \text{H}$ (Example 148)

243e : $n = 2$, $\text{R}_1 = \text{NO}_2$, $\text{R}_2 = \text{H}$ (Example 149)

243f : $n = 2$, $\text{R}_1 = \text{R}_2 = \text{Cl}$ (Example 150)

243g : $n = 2$, $\text{R}_1 = \text{OH}$, $\text{R}_2 = \text{H}$ (Example 151)

243h : $n = 2$, $\text{R}_1 = \text{OH}$, $\text{R}_2 =$  (Example 152)

Example 145:

5 N-(2-Aminophenyl)-6-(N-phthaloyl)hexanamide (243a)

Step 1: 6-N-Phthaloylhexanoic acid (242a)

To a solution of phthalic anhydride 241a (1.48 g, 10.00 mmol) in anhydrous ethyl alcohol (20 mL) was added 6-aminocaproic acid (1.97 g, 15.00 mmol). The

reaction mixture was refluxed for 16 hours, cooled to room temperature, evaporated to dryness to produce a solid material, which was triturated with water to form a suspension. The suspension was filtered, the solid was collected and dried to afford the title compound 242a (1.18 g, 45% yield). ¹H NMR: (400 MHz, DMSO-d₆) δ 7.87-7.80 (m, 4H), 3.54 (t, J = 7.1 Hz, 2H), 2.17 (t, J = 7.3 Hz, 2H), 1.62-1.44 (m, 4H), 1.30-1.23 (m, 2H).

Step 2: N-(2-Aminophenyl)-6-(N-phthaloyl)hexanamide (243a)

Following the procedure described in Example 22, but substituting carboxylic acid 242a for 50g, the title compound 243a was obtained in 31% yield. ¹H NMR: (400 MHz, DMSO-d₆) δ 9.05 (s, 1H), 7.88-7.80 (m, 4H), 7.09 (dd, J = 7.8, 1.4 Hz, 1H), 6.86 (dd, J = 8.0, 1.5 Hz, 1H), 6.68 (dd, J = 8.0, 1.4 Hz, 1H), 6.49 (dd, J = 7.8, 1.5 Hz, 1H), 4.80 (s, 2H), 3.57 (t, J = 7.0 Hz, 2H), 2.28 (t, J = 7.3 Hz, 2H), 1.64-1.57 (m, 4H), 1.37-1.27 (m, 2H),

Example 146:

N-(2-Aminophenyl)-5-(N-phthaloyl)pentanamide (243b)

Step 1: 5-(N-Phthaloyl)pentanoic acid (242b)

Following a procedure analogous to that described in Example 145, step 1, but substituting 5-amino-valeric acid for 6-amino-caproic acid, the title compound 242b was obtained in 46% yield. MS (ESI) = 270 (M+Na⁺).

Step 2: N-(2-Aminophenyl)-5-(N-phthaloyl)-pentanamide (243b)

Following the procedure described in Example 22, but substituting carboxylic acid 242b for 50g, the title compound 243b was obtained in 28% yield. ¹H NMR: (300 MHz, DMSO-d₆) δ 9.06 (s, 1H), 7.87-7.80 (m, 4H), 7.11 (d, J = 7.7 Hz, 1H), 6.86 (t, J = 7.4 Hz, 1H), 6.68 (dd, J = 8.0, 1.4 Hz, 1H), 6.50 (dd, J = 8.8, 1.1 Hz, 1H), 4.78 (br s, 2H), 3.59 (t, J = 5.8 Hz, 2H), 2.33 (t, J = 6.6 Hz, 2H), 1.60 (m, 4H).

Example 147:***N*-(2-Aminophenyl)-7-(*N*-phthaloyl)heptanamide (243c)****Step 1: 7-(*N*-Phthaloyl)heptanoic acid (242c)**

Following a procedure analogous to that described in Example 145, step 1,
5 but substituting 7-aminoheptanoic acid for 6-amino-caproic acid, the title compound 242c was obtained in 46% yield. MS (ESI) = 298 (MH⁺).

Step 2: *N*-(2-Aminophenyl)-7-(*N*-phthaloyl)-heptanamide (243c)

Following the procedure described in Example 22, but substituting
carboxylic acid 242c for 50g, the title compound 243c was obtained in 28% yield.
10 ¹H NMR: (300 MHz, DMSO-d₆) δ 9.05 (s, 1H), 7.87-7.79 (m, 4H), 7.11 (d, J = 7.7 Hz, 1H), 6.86 (t, J = 7.4 Hz, 1H), 6.68 (d, J = 8.0 Hz, 1H), 6.50 (t, J = 7.7 Hz, 1H), 4.77 (s, 2H), 3.55 (t, J = 7.1 Hz, 2H), 2.27 (t, J = 7.4 Hz, 2H), 1.61-1.53 (m, 4H), 1.31-1.28 (m, 4H).

Example 148:***N*-(2-Aminophenyl)-6-(5-methyl-*N*-phthaloyl)hexanamide (243d)****Step 1: 6-(5-Methyl-*N*-phthaloyl)hexanoic acid (242d)**

Following a procedure analogous to that described in Example 145, step 1,
but substituting 5-methyl-*N*-phthaloyl for *N*-phthaloyl, the title compound 242d
was obtained in 85% yield. MS (ESI) = 276 (MH⁺).

Step 2: *N*-(2-Aminophenyl)-6-(5-methyl-*N*-phthaloyl)hexanamide (243d)

Following the procedure described in Example 22, but substituting
carboxylic acid 242d for 50g, the title compound 243d was obtained in 43% yield.
¹H NMR: (300 MHz, DMSO-d₆) δ 9.04 (s, 1H), 7.73-7.60 (m, 3H), 7.07 (d, J = 7.7 Hz, 1H), 6.85 (t, J = 8.0 Hz, 1H), 6.67 (dd, J = 8.0, 1.1 Hz, 1H), 6.50 (t, J = 7.7 Hz, 1H),
25 4.77 (s, 2H), 3.52 (t, J = 7.1 Hz, 2H), 2.46 (s, 3H, CH₃), 2.27 (t, J = 7.1 Hz, 2H), 1.58 (br s, 4H), 1.32-1.22 (m, 2H).

Example 149:***N*-(2-Aminophenyl)-6-(5-nitro-*N*-phthaloyl)hexanamide (243e)****Step 1: 6-(5-Nitro-*N*-phthaloyl)hexanoic acid (242e)**

- Following a procedure analogous to that described in Example 145, step 1, but substituting 5-nitro-*N*-phthaloyl for *N*-phthaloyl, the title compound 242e was obtained in 26% yield. MS (ESI) = 305 (MH⁺).

Step 2: *N*-(2-Aminophenyl)-6-(5-nitro-*N*-phthaloyl)hexanamide (243e)

- Following the procedure described in Example 22, but substituting carboxylic acid 242e for 50g, the title compound 243e was obtained in 14% yield.
- ¹H NMR: (400 MHz, DMSO-d₆) δ 9.05 (s, 1H), 8.60 (dd, J = 8.2 Hz, 1.9 Hz, 1H), 8.47 (d, J = 1.5 Hz, 1H), 8.10 (d, J = 8.2 Hz, 1H), 7.10 (dd, J = 7.9, 1.3 Hz, 1H), 6.86 (dd, J = 8.0, 1.5 Hz, 1H), 6.67 (dd, J = 1.3, J = 8.0 Hz, 1H), 6.49 (dd, J = 7.7, 1.4 Hz, 1H), 4.76 (s, 2H), 3.62 (t, J = 7.0 Hz, 2H), 2.29 (t, J = 7.3 Hz, 2H), 1.71-1.55 (m, 4H), 1.38-1.30 (m, 2H).

Example 150:***N*-(2-Aminophenyl)-6-(5,6-dichloro-*N*-phthaloyl)hexanamide (243f)****Step 1: 6-(5,6-Dichloro-*N*-phthaloyl)hexanoic acid (242f)**

- Following a procedure analogous to that described in Example 145, step 1, but substituting 5,6-dichloro-*N*-phthaloyl for *N*-phthaloyl, the title compound 242f was obtained in 64% yield. MS (ESI) = 328 (MH⁺).

Step 2: *N*-(2-Aminophenyl)-6-(5,6-dichloro-*N*-phthaloyl)hexanamide (243f)

- Following the procedure described in Example 22, but substituting carboxylic acid 242f for 50g, the title compound 243f was obtained in 52% yield.
- ¹H NMR: (300 MHz, DMSO-d₆) δ 9.04 (s, 1H), 8.15 (s, 2H), 7.07 (d, J = 7.4 Hz, 1H), 6.85 (t, J = 7.8 Hz, 1H), 6.67 (d, J = 8.0 Hz, 1H), 6.48 (t, J = 7.7 Hz, 1H), 4.76 (s, 2H), 3.56 (t, J = 6.9 Hz, 2H), 2.27 (t, J = 7.1 Hz, 2H), 1.63-1.56 (m, 4H), 1.34-1.27 (m, 2H).

Example 151:***N*-(2-Aminophenyl)-6-(5-hydroxy-*N*-phthaloyl)hexanamide (243g)****Step 1: 6-(5-Hydroxy-*N*-phthaloyl)hexanoic acid (242g)**

4-Hydroxy-phthalic acid 240 (2.30 g, 12.64 mmol) was heated for two hours at 220°C under vacuum, cooled to room temperature and dissolved in 50 mL anhydrous ethyl alcohol. To this solution 6-amino caproic acid (2.48 g, 18.93 mmol) was added and the mixture was refluxed for 28 hours, cooled and evaporated to produce an oily residue. This material was triturated with water to form a precipitate which was collected and dried to afford the title compound 242g (2.24 g, 64% yield). MS (ESI) = 276 (MH⁺).

Step 2: *N*-(2-Aminophenyl)-6-(5-hydroxy-*N*-phthaloyl)hexanamide (243g)

Following the procedure described in Example 22, but substituting carboxylic acid 242g for 50g, the title compound 243g was obtained in 11% yield. ¹H NMR: (300 MHz, DMSO-d₆) δ 10.88 (br s, 1H), 9.04 (s, 1H), 7.71 (d, J = 8.0 Hz, 4H), 7.11-7.07 (m, 3H), 6.86 (d, J = 6.3 Hz, 1H), 6.68 (d, J = 6.3 Hz, 1H), 6.49 (t, J = 7.4 Hz, 1H), 4.77 (s, 2H), 3.51 (t, J = 7.1 Hz, 2H), 2.27 (t, J = 7.4 Hz, 2H), 1.62-1.54 (m, 4H), 1.37-1.25 (m, 2H).

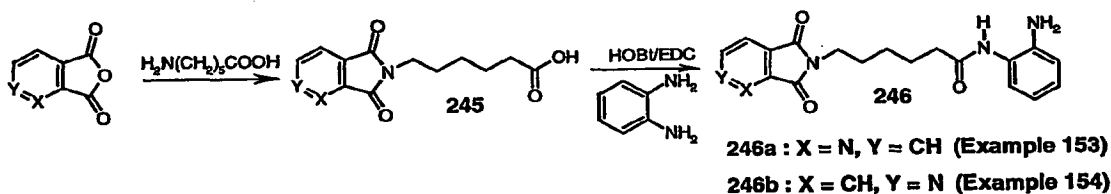
Example 152:***N*-(2-Aminophenyl)-6-(5-hydroxy-6-(1-methylpiperidinyl)-*N*-phthaloyl)-hexanamide (243h)****Step 1: 6-(5-Hydroxy-6-(1-methylpiperidinyl)-*N*-phthaloyl)hexanoic acid (244)**

To a stirring suspension of the phthalimido acid 242g (500 mg, 1.81 mmol) in a mixture of anhydrous dioxane (7 mL) and anhydrous DMF (3.5 mL) at room temperature was added paraformaldehyde (73 mg, 2.4 mmol) followed by addition of piperidine (0.22 mL, 2.2 mmol). The reaction mixture was stirred at 80°C for 8 hours, solvents were removed in vacuum and the crude residue was purified by column chromatography using a gradient of 100% dichloromethane to 5% methanol in dichloromethane. The resulting yellow solid was crystallized

from methanol-diethyl ether to afford the title compound 244 (0.25 g, 37%). MS (ESI) = 375 (MH⁺).

Step 2: *N*-(2-Aminophenyl)-6-(5-hydroxy-6-(1-methylpiperidinyl)-*N*-phthaloyl)hexanamide (243h)

- 5 Following the procedure described in Example 22, but substituting carboxylic acid 244 for 50g, the title compound 243h was obtained in 44% yield. ¹H NMR: (300 MHz, DMSO-d₆) δ 9.04 (s, 1H), 7.58 (s, 1H), 7.07 (d, J = 7.97 Hz, 1H), 7.02 (s, 1H), 6.85 (t, J = 7.69 Hz, 1H), 6.67 (d, J = 7.97 Hz, 1H), 6.48 (t, J = 7.69 Hz, 1H), 4.76 (br s, 2H), 3.75 (s, 2H), 3.50 (t, J = 6.87 Hz, 2H), 2.26 (t, J = 7.28 Hz, 2H), 1.54 (m, 8H), 1.44 (m, 2H), 1.27 (m, 2H).
- 10



Example 153:

- 15 ***N*-(2-Aminophenyl)-6-(5,7-dioxo-5,7-dihydro-pyrrolo[3,4-*b*]pyridinyl)hexanamide (246a)**

Step 1: 5-[6-(5,7-Dioxo-5,7-dihydro-pyrrolo[3,4-*b*]pyridinyl)]hexanoic acid (245a)

Following a procedure analogous to that described in Example 145, step 1, but substituting 2,3-pyridinedicarboxylic anhydride for *N*-phthalic anhydride, the title compound 245a was obtained in 22% yield. MS (ESI) = 363 (MH⁺).

- 20 **Step 2: *N*-(2-Aminophenyl)-6-[6-(5,7-dioxo-5,7-dihydro-pyrrolo[3,4-*b*]pyridinyl)]hexanamide (246a)**

Following the procedure described in Example 22, but substituting carboxylic acid 245a for 50g, the title compound 246a was obtained in 40% yield. ¹H NMR: (300 MHz, DMSO-d₆) δ 9.06 (s, 1H), 8.94 (dd, J = 4.9, 1.4 Hz, 1H), 8.27 (dd, J = 7.7, 1.4 Hz, 1H), 7.75 (dd, J = 7.7, 4.9 Hz, 1H), 7.08 (d, J = 8.0 Hz, 1H), 6.85

25

(t, J = 8.2 Hz, 1H), 6.67 (d, J = 8.2 Hz, 1H), 6.49 (t, J = 7.7 Hz, 1H), 4.76 (s, 2H), 3.60 (t, J = 7.1 Hz, 2H), 2.28 (t, J = 6.9 Hz, 2H), 1.63-1.57 (m, 4H), 1.35-1.28 (m, 2H).

Example 154:

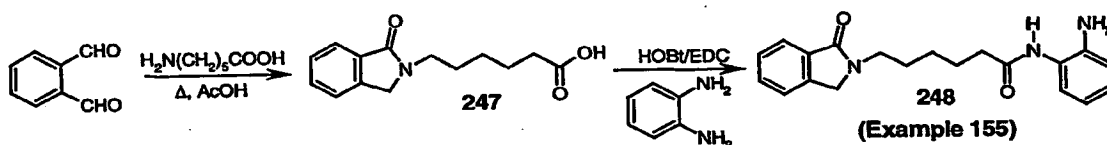
N-(2-Aminophenyl)-6-(5,7-dioxo-5,7-dihydro-pyrrolo[3,4-c]pyridinyl)]hexanamide (246b)

Step 1: 5-[6-(5,7-Dioxo-5,7-dihydro-pyrrolo[3,4-c]pyridinyl)]hexanoic acid (245b)

Following a procedure analogous to that described in Example 145, step 1, but substituting 3,4-pyridinedicarboxylic anhydride for N-phthalic anhydride, the title compound 245b was obtained in 8% yield. MS (ESI) = 363 (MH⁺).

Step 2: N-(2-Aminophenyl)-6-[6-(5,7-Dioxo-5,7-dihydro-pyrrolo[3,4-c]pyridinyl)]hexanamide (246b)

Following the procedure described in Example 22, but substituting carboxylic acid 245b for 50g, the title compound 246b was obtained in 61% yield. ¹H NMR: (300 MHz, DMSO-d₆) δ 9.05-9.09 (m, 3H), 7.87 (d, J = 4.7 Hz, 1H), 7.08 (d, J = 7.4 Hz, 1H), 6.86 (t, J = 7.1 Hz, 1H), 6.67 (d, J = 6.9 Hz, 1H), 6.49 (t, J = 6.3 Hz, 1H), 4.77 (s, 2H), 3.58 (t, J = 6.9 Hz, 2H), 2.28 (t, J = 7.1 Hz, 2H), 2.00-1.90 (m, 2H), 1.64-1.54 (m, 4H), 1.33-1.29 (m, 2H).



Example 155:

N-(2-Aminophenyl)-6-[2-(1-oxo-1,3-dihydro-isoindolyl)]hexanamide (248)

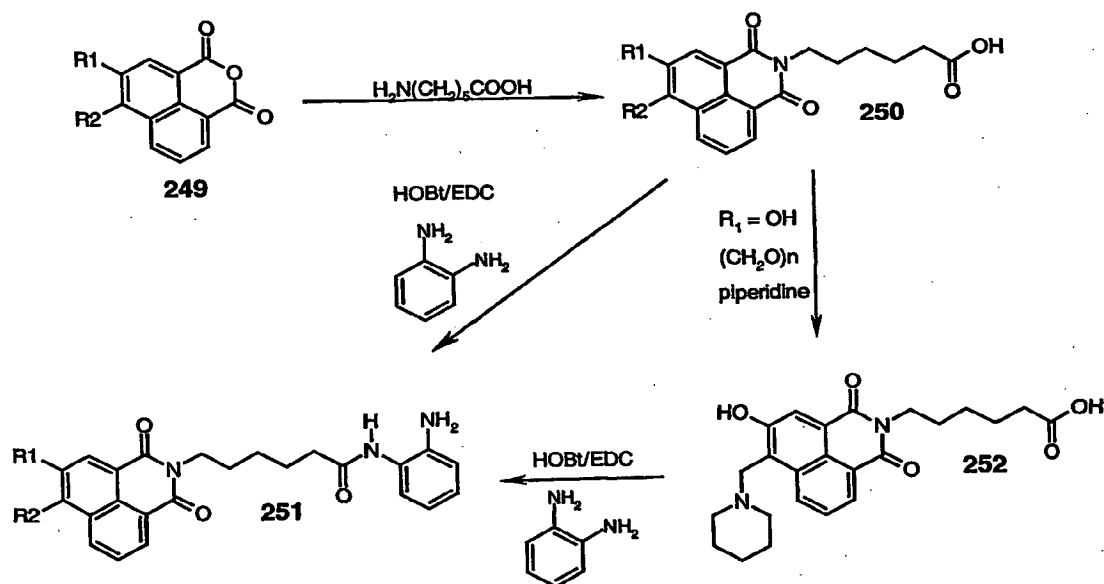
Step 1: 6-[2-(1-Oxo-1,3-dihydro-isoindolyl)]hexanoic acid (247)

To a solution of benzene-1,2-carbaldehyde (670 mg, 5.00 mmol) in acetic acid (6 mL) was added 6-amino-hexanoic acid (655 mg, 5.00 mmol). The reaction mixture was refluxed for 15 minutes, cooled to room temperature and evaporated.

The residue was chromatographed on a silica gel column (5% to 10% methanol in dichloromethane) to produce the title compound 247 (1.20 g, 97% yield). MS (ESI) = 248 (MH⁺).


Step 2: *N*-(2-Aminophenyl)-6-[2-(1-oxo-1,3-dihydro-isoindolyl)]hexanamide (248)

- 5 Following the procedure described in Example 22, but substituting
 carboxylic acid 247 for 50g, the title compound 248 was obtained in 27% yield. ¹H
 NMR: (300 MHz, DMSO-d₆) δ 9.06 (s, 1H), 7.66-7.46 (m, 4H), 7.09 (d, J = 8.0 Hz,
 1H), 6.86 (t, J = 7.7 Hz, 1H), 6.68 (d, J = 7.7 Hz, 1H), 6.49 (t, J = 7.7 Hz, 1H), 4.78 (s,
 2H), 4.45 (s, 2H), 3.51 (t, J = 6.9 Hz, 2H), 2.29 (t, J = 7.1 Hz, 2H), 1.63-1.60 (m, 4H),
 10 1.33-1.27 (m, 2H).



251a : R1 = OH, R2 = H (Example 156)

251b : R1 = H, R2 = Br (Example 157)

251c : R1 = OH, R2 =  **(Example 158)**

Example 156:

***N*-(2-Aminophenyl)-6-[2-(5-hydroxy-1,3-dioxo-1*H*,3*H*-benzo[*de*]isoquinoliny)]-hexanamide (251a)**

5 **Step 1: 6-[2-(5-Hydroxy-1,3-dioxo-1*H*,3*H*-benzo[*de*]isoquinoliny)]hexanoic acid (250a)**

Following a procedure analogous to that described in Example 145, step 1, but substituting 3-hydroxy-1,8-naphthalic anhydride for *N*-phthalic anhydride, the title compound 250a was obtained in 98% yield. MS (ESI) = 328 (MH⁺).

10 **Step 2: *N*-(2-Aminophenyl)-6-[2-(5-hydroxy-1,3-dioxo-1*H*,3*H*-benzo[*de*]isoquinoliny)]- hexanamide (251a)**

Following the procedure described in Example 22, but substituting carboxylic acid 250a for 50g, the title compound 251a was obtained in 18% yield. ¹H NMR: (400 MHz, DMSO-*d*₆) δ 10.52 (s, 1H), 9.07 (s, 1H), 8.22 (t, *J* = 6.5 Hz, 2H), 8.00 (d, *J* = 2.4 Hz, 1H), 7.72 (t, *J* = 7.4 Hz, 1H), 7.63 (d, *J* = 2.4 Hz, 1H), 7.09 (dd, *J* = 7.8 Hz, 1.3 Hz, 1H), 6.87 (dd, *J* = 8.0 Hz, 1.4 Hz, 1H), 6.69 (dd, *J* = 8.0 Hz, 1.3 Hz, 1H), 6.49 (dd, *J* = 7.7 Hz, 1.4 Hz, 1H), 4.81 (s, 2H), 4.02 (t, *J* = 7.3 Hz, 2H), 2.31 (t, *J* = 7.3 Hz, 2H), 1.67-1.58 (m, 4H), 1.42-1.35 (m, 2H).

Example 157:

20 ***N*-(2-Aminophenyl)-6-[2-(6-bromo-1,3-dioxo-1*H*,3*H*-benzo[*de*]isoquinoliny)]-hexanamide (251b)**

Step 1: 6-[2-(6-Bromo-1,3-dioxo-1*H*,3*H*-benzo[*de*]isoquinoliny)]hexanoic acid (250b)

25 Following a procedure analogous to that described in Example 145, step 1, but substituting 4-bromo-1,8-naphthalic anhydride for *N*-phthalic anhydride, the title compound 250a was obtained in 96% yield. MS (ESI) = 390 (MH⁺).

Step 2: *N*-(2-Aminophenyl)-6-[2-(6-bromo-1,3-dioxo-1*H*,3*H*-benzo[*de*]isoquinoliny)]- hexanamide (251b)

Following the procedure described in Example 22, but substituting carboxylic acid 250b for 50g, the title compound 251b was obtained in 26% yield.

¹H NMR: (300 MHz, DMSO-d₆) δ 9.05 (s, 1H, NH), 8.52 (t, J = 8.2 Hz, 2H), 8.30 (d, J = 7.7 Hz, 1H), 8.18 (d, J = 7.7 Hz, 1H), 7.96 (t, J = 7.7 Hz, 1H), 7.06 (d, J = 8.0 Hz, 1H), 6.85 (t, J = 8.0 Hz, 1H), 6.67 (d, J = 8.0 Hz, 1H), 6.47 (t, J = 7.7 Hz, 1H), 4.78 (br s, 2H), 4.02 (t, J = 7.1 Hz, 2H), 2.30 (t, J = 7.1 Hz, 2H), 1.66-1.58 (m, 4H), 1.41-1.34 (m, 2H).

Example 158:

N-(2-Aminophenyl)-6-[2-(5-hydroxy-1,3-dioxo-6-(1-methylpiperidynyl)-1*H*,3*H*-benzo[*de*]isoquinoliny)]-hexanamide (251c)

10 Step 1: 6-[2-(5-Hydroxy-1,3-dioxo-6-piperidyn-1-ylmethyl-1*H*,3*H*-benzo[*de*]isoquinoliny)]hexanoic acid (252)

To a stirring suspension of the acid 250a (390 mg, 1.19 mmol) in a mixture of anhydrous dioxane (5 mL) and anhydrous DMF (2 mL) at room temperature was added paraformaldehyde (70 mg, 2.33 mmol) followed by addition of piperidine (0.14 mL, 1.42 mmol). The reaction mixture was stirred for 18 hours at ambient temperature, solvents were removed in vacuum and the crude residue was triturated with 0.5 mL methanol followed by addition of 25 mL ether. The formed suspension was filtered to afford the title compound 252 as yellow crystals (424 mg, 84%). MS (ESI) = 425 (MH⁺).

20 Step 2: *N*-(2-Aminophenyl)-6-[2-(5-hydroxy-1,3-dioxo-6-(1-methylpiperidynyl)-1*H*,3*H*-benzo[*de*]isoquinoliny)]hexanamide (251c)

Following the procedure described in Example 22, but substituting carboxylic acid 252 for 50g, the title compound 251c was obtained in 39% yield. ¹H NMR: (300 MHz, DMSO-d₆) δ 9.05 (s, 1H), 8.43 (d, J = 8.2 Hz, 1H), 8.27 (d, J = 7.1 Hz, 1H), 7.96 (s, 1H), 7.75 (t, J = 8.2 Hz, 1H), 7.08 (d, J = 8.0 Hz, 1H), 6.85 (t, J = 6.7 Hz, 1H), 6.68 (d, J = 8.0 Hz, 1H), 6.47 (t, J = 7.1 Hz, 1H), 4.77 (s, 2H), 4.15 (s, 2H), 4.02 (t, J = 6.9 Hz, 2H), 2.54 (br s, 4H), 2.30 (t, J = 7.1 Hz, 2H), 1.65-1.36 (m, 12H).

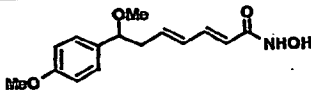
Example 159:**Inhibition of Histone Deacetylase Enzymatic Activity**

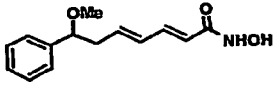
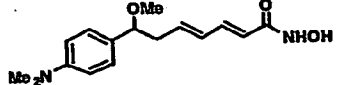
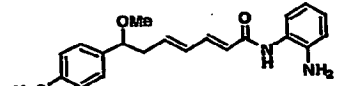
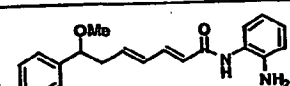
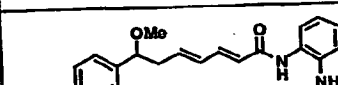
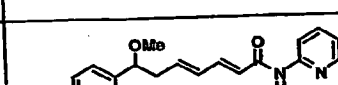
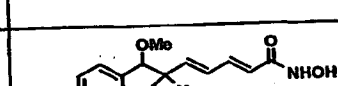
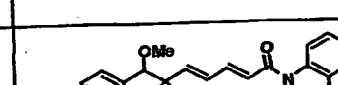
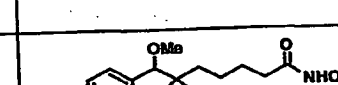
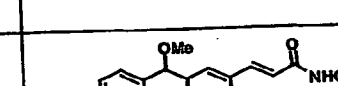
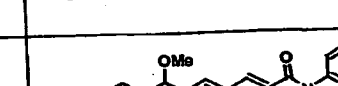
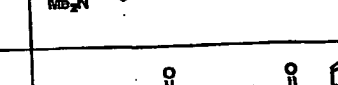
HDAC inhibitors were screened against histone deacetylase enzyme in nuclear extracts prepared from the human small cell lung cancer cell line H446 (ATTC HTB-171) and against a cloned recombinant human HDAC-1 enzyme expressed and purified from a Baculovirus insect cell expression system.

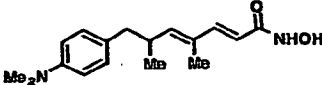
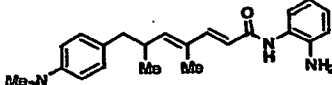
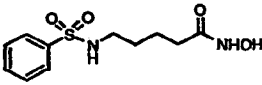
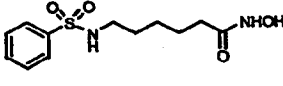
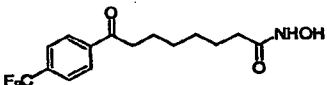
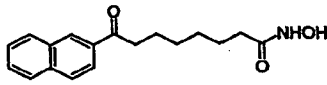
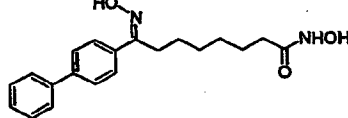
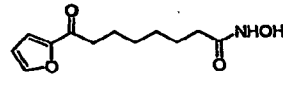
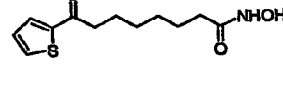
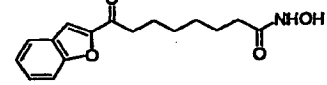
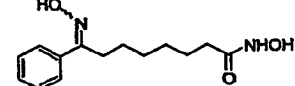
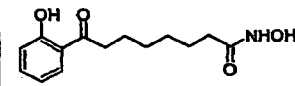
For deacetylase assays, 20,000 cpm of the [³H]-metabolically labeled acetylated histone substrate (M. Yoshida *et al.*, *J. Biol. Chem.* 265(28): 17174-17179 (1990)) was incubated with 30 µg of H446 nuclear extract or an equivalent amount of the cloned recombinant hHDAC-1 for 10 minutes at 37 °C. The reaction was stopped by adding acetic acid (0.04 M, final concentration) and HCl (250 mM, final concentration). The mixture was extracted with ethyl acetate and the released [³H]-acetic acid was quantified by scintillation counting. For inhibition studies, the enzyme was preincubated with compounds at 4 °C for 30 minutes prior to initiation of the enzymatic assay. IC₅₀ values for HDAC enzyme inhibitors were determined by performing dose response curves with individual compounds and determining the concentration of inhibitor producing fifty percent of the maximal inhibition.

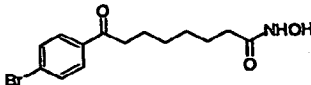
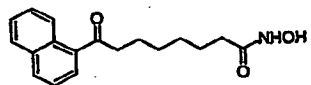
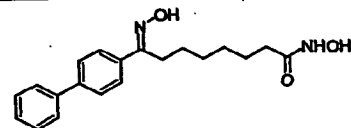
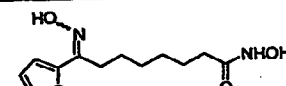
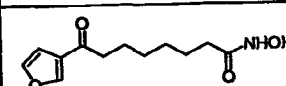
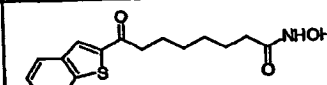
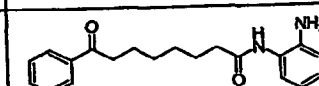
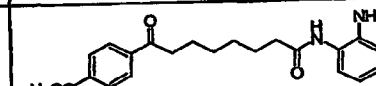
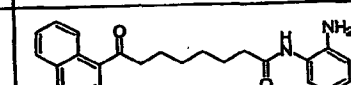
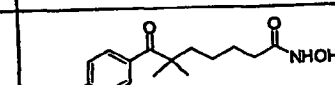
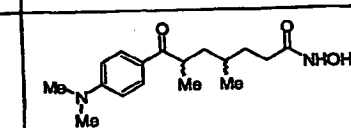
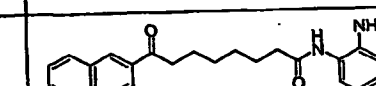
Representative data are presented in Table 2. In the first column are reported IC₅₀ values determined against histone deacetylase in nuclear extracts from H446 cells (pooled HDACs). In the second column are reported IC₅₀ values determined against recombinant human HDAC-1 enzyme (rHDAC-1).

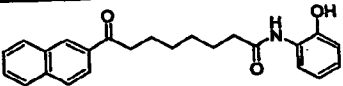
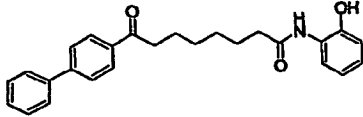
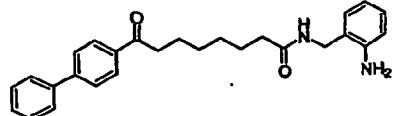
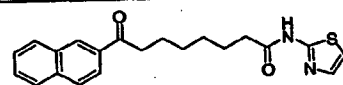
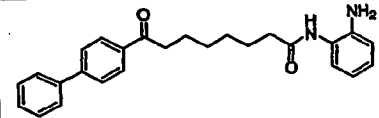
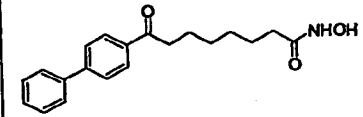
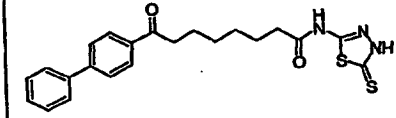
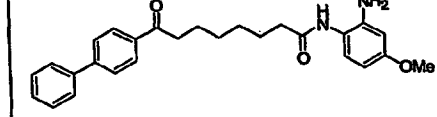
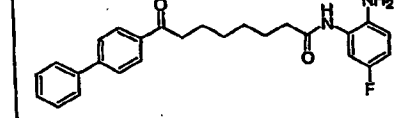
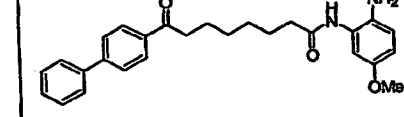
Table 2: Inhibition of Histone Deacetylase

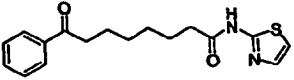
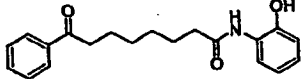
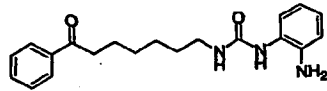
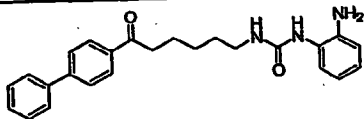
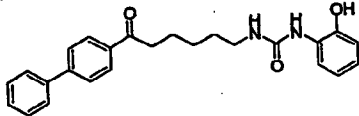
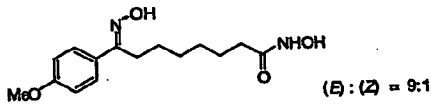
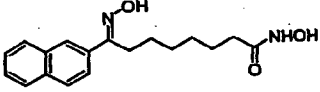
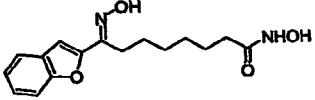
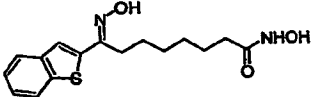
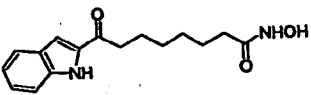
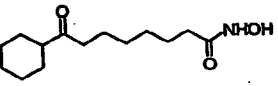
Example	Cpd.	Structure	pooled HDACs IC ₅₀ (µM)	rHDAC-1 IC ₅₀ (µM)
Ex. 1	6		3	0.25

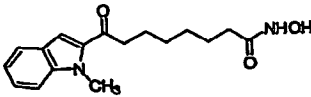
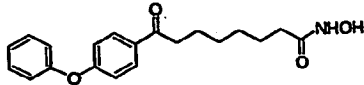
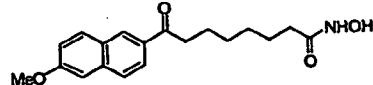
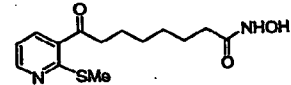
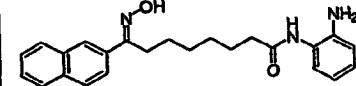
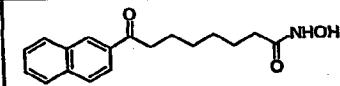
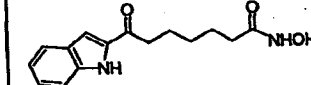
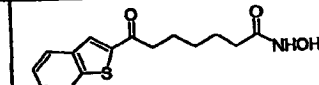
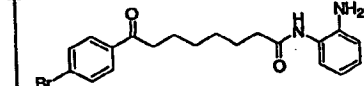
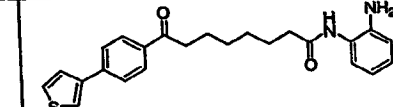
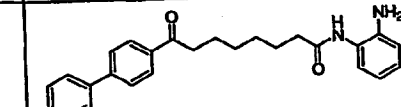
Ex. 2	12		9.4	ND
Ex. 3	17		4	0.25
Ex. 4	18		> 100	1
Ex. 5	19		> 100	<20
Ex. 6	20		> 20	3
Ex. 7	21		> 100	ND
Ex. 8	26		5	1
Ex. 9	27		> 100	ND
Ex. 10	28		4	0.1
Ex. 11	33		> 10	ND
Ex. 12	34		> 100	7
Ex. 13	35		> 100	2

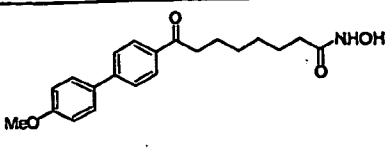
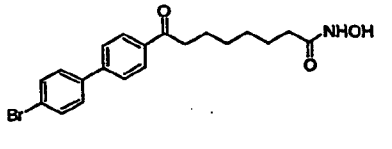
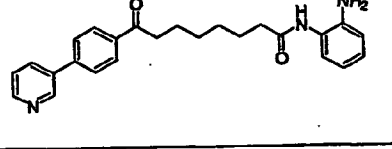
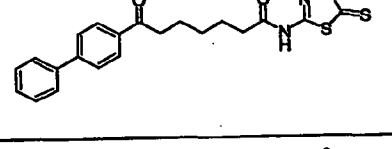
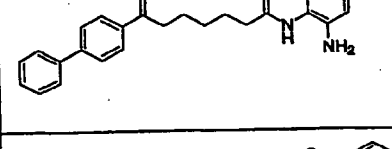
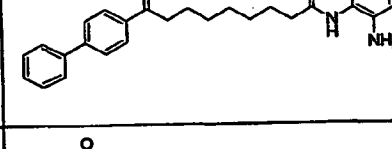
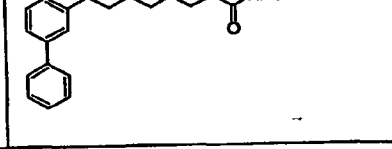
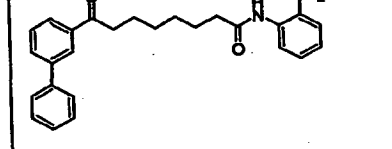
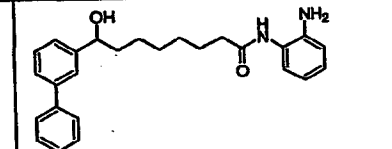
Ex. 14	38		> 20	ND
Ex. 15	39		> 20	7
Ex. 16	42		6	1
Ex. 17	47		4.5	0.5
Ex. 18	52c		0.55	0.06
Ex. 18	52g		0.03	0.005
Ex. 18	(Z)-51h		75%	
Ex. 18	52i		3	0.1
Ex. 18	52k		0.9	0.18
Ex. 18	52l		0.3	0.03
Ex. 18	51a		0.35	0.1
Ex. 18	52e		0.55	0.06

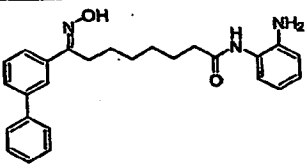
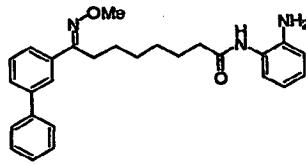
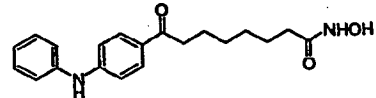
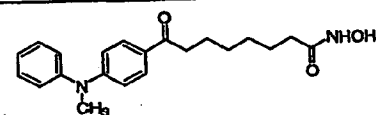
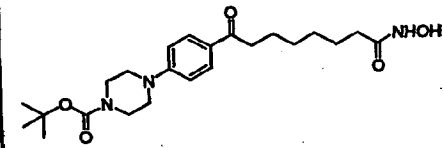
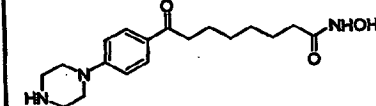
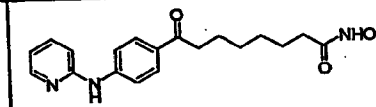
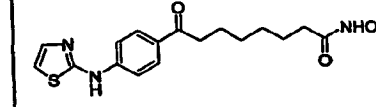
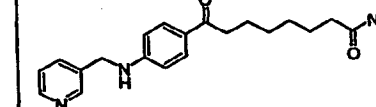
Ex. 18	52d		0.05	0.055
Ex. 18	52f		0.55	0.01
Ex. 18	(E)-51h		99%	
Ex. 18	51i		0.9	0.1
Ex. 18	52j		3	0.45
Ex. 18	52m		0.1	0.01
Ex. 19	53a			0.6
Ex. 19	53b		0%	1
Ex. 19	53f		3%	2
Ex. 20	58		3	0.25
Ex. 21	62		1.1	
Ex. 22	63		> 20	2.2

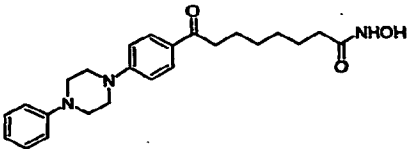
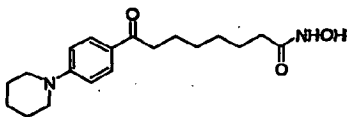
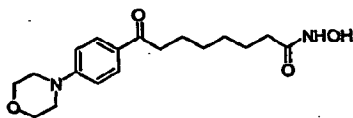
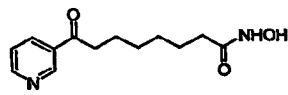
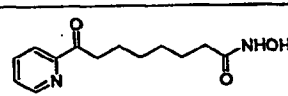
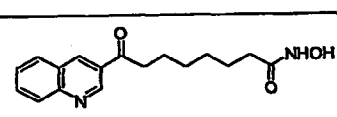
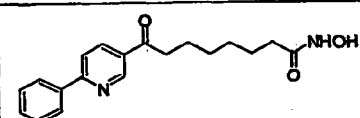
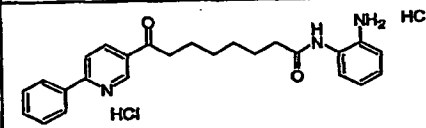
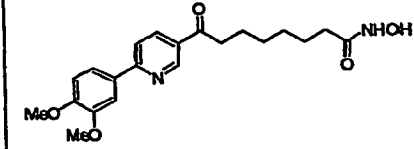
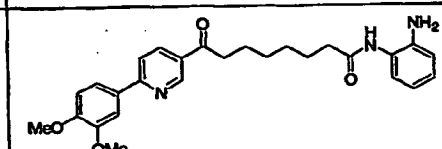
Ex. 25	66		0%	2.3
Ex. 29	74		0%	7.8
Ex. 30	75		0%	8.4
Ex. 31	76			23
Ex. 32	77		11%	1.9
Ex. 33	78		0.04	0.004
Ex. 34	79			9
Ex. 36	81		27%	25
Ex. 37	82			20
Ex. 38	83			25

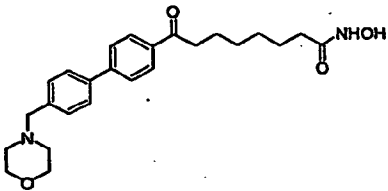
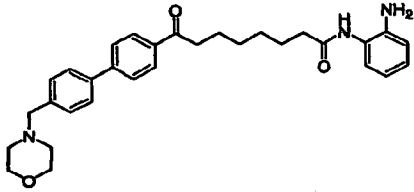
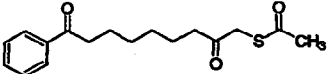
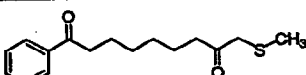
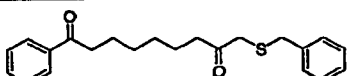
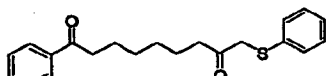
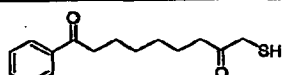
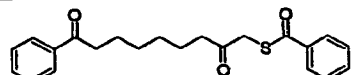


Ex. 40	85			20
Ex. 41	86			5
Ex. 43	89		0%	16.3
Ex. 44	92		0%	15.9
Ex. 45	93		0%	6.4
Ex. 48	96		0.1	0.035
Ex. 49	97		0.03	0.01
Ex. 50	98		0.04	0.03
Ex. 51	99		0.035	0.008
Ex. 52	100			4
Ex. 54	102			

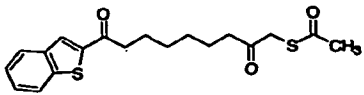
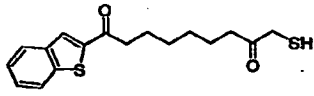
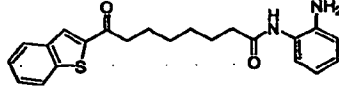
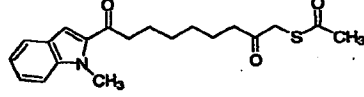
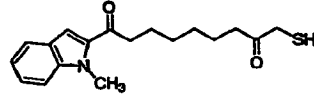
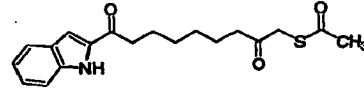
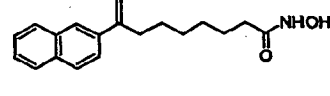
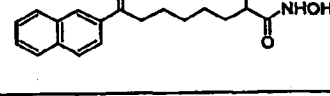
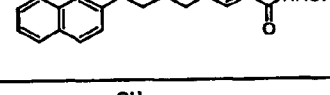
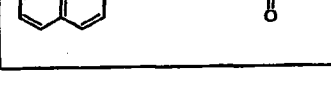
Ex. 55	103		0.1	0.05
Ex. 56	104		61% @ 1 μ M	0.16
Ex. 57a	105a		83% @ 1 μ M	0.01
Ex. 57b	105b		40% @ 1 μ M	0.02
Ex. 58	106		> 20	4
Ex. 63	111		0.35	0.2
Ex. 64	112		0.25	0.039
Ex. 65	113		33% @ 1 μ M	0.13
Ex. 66	114		> 20	6.3
Ex. 67	117		0%	9
Ex. 68	123			4

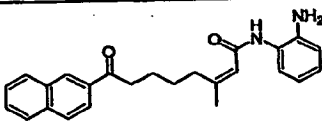
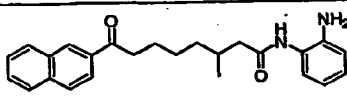
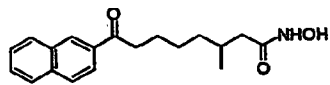
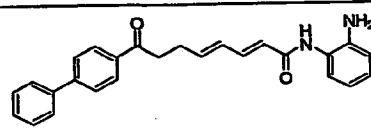
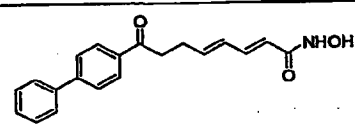
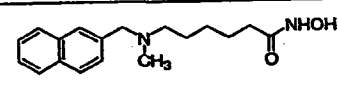
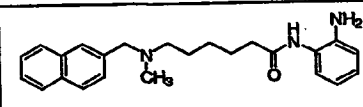
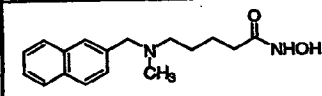
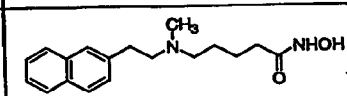
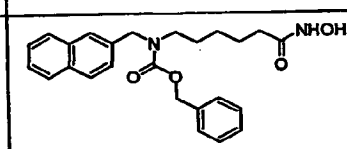
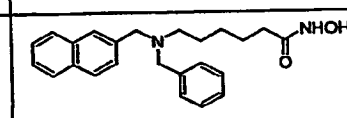
Ex. 69	120		83%	0.002
Ex. 70	124			0.001
Ex. 71	127		0%	10.3
Ex. 72	128			19
Ex. 73	129			1
Ex. 74	130			3
Ex. 75	131			0.008
Ex. 76	132			0.8
Ex. 77	133			1

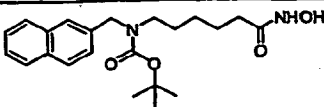
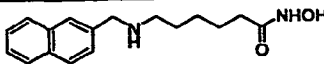
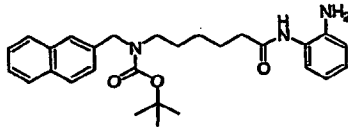
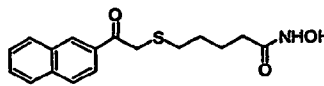
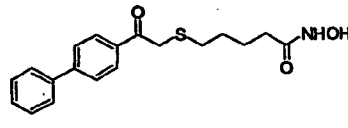
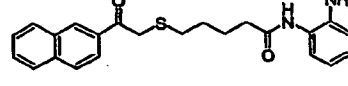
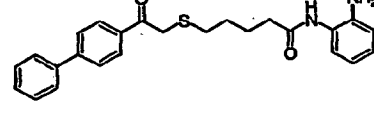
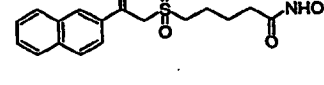
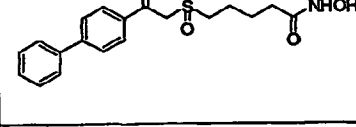
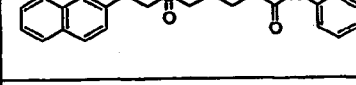
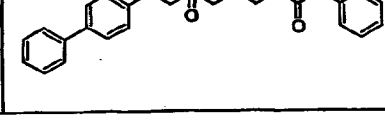
Ex. 78	134			1
Ex. 79	135			2
Ex. 80	138a		83% @ 1 μ M	0.004
Ex. 81	138b		0.049	0.008
Ex. 82a	138c1			0.03
Ex. 82b	138c2			0.03
Ex. 83	138d		0.02	0.004
Ex. 84	138e		0.02	0.004
Ex. 85	138f			0.02

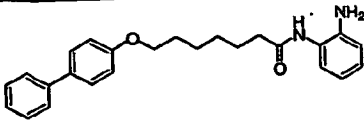
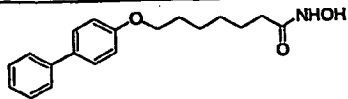
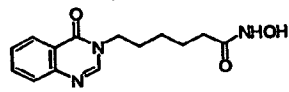
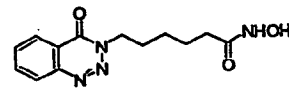
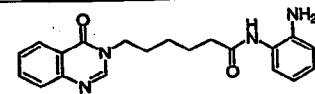
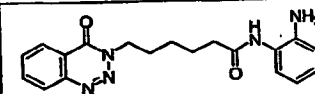
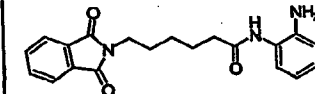
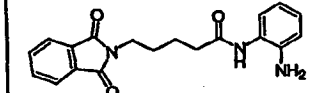
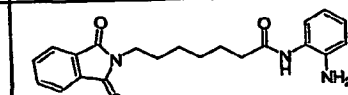
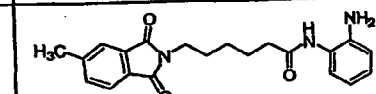
Ex. 86	138g		0.051	0.006
Ex. 87	138h		0.048	0.009
Ex. 88	138i		0.1	0.033
Ex. 89a	141a1		22% @ 1 μ M	0.06
Ex. 89b	141a2		12% @ 1 μ M	
Ex. 90	141b		71% @ 1 μ M	0.035
Ex. 91	145			0.008
Ex. 92	146			2
Ex. 93	150			0.05
Ex. 94	151			4

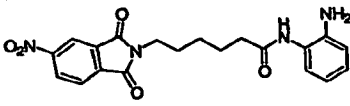
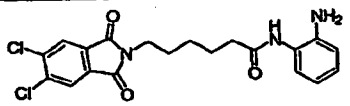
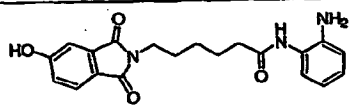
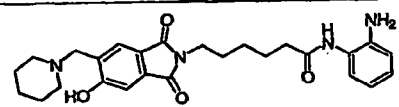
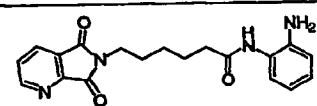
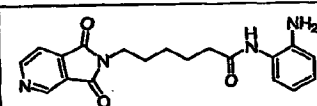
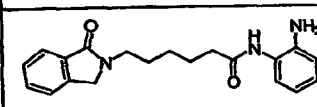
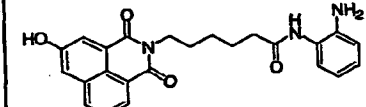
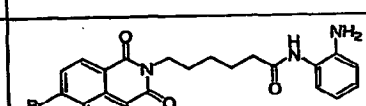
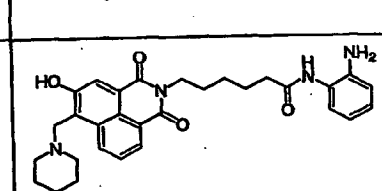
Ex. 95	156			20
Ex. 96	157			4
Ex. 97	159a		0.05	0.1
Ex. 98	159b		14% @ 5 μ M	0.5
Ex. 99	159c		0% @ 5 μ M	5
Ex. 100	159d		50	5
Ex. 101	159e		0.05	0.05
Ex. 102	159f		0.35	0.7
Ex. 103	160		0.08	0.13
Ex. 104	161		0.094	0.071

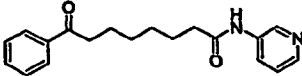
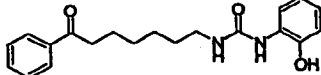
Ex. 105	162		0.04	0.035
Ex. 106	163		0.04	0.01
Ex. 107	164		18%	2.3
Ex. 108	165		0.04	0.09
Ex. 109	166		0.05	0.42
Ex. 110	167		0.3	0.4
Ex. 111	170			0.01
Ex. 112	175		2.5	0.5
Ex. 115	187			0.04
Ex. 116	188			0.02

Ex. 117	194			24
Ex. 118	198			16
Ex. 119	199		0.08	0.03
Ex. 120	204			5
Ex. 121	205			0.04
Ex. 122	209			0.05
Ex. 123	210			10
Ex. 124	213			3
Ex. 125	216			0.3
Ex. 126	220a			0.02
Ex. 127	220b			0.1

Ex. 128	222			0.1
Ex. 129	223			2
Ex. 130	224			3
Ex. 131	226a			0.007
Ex. 132	226b			0.006
Ex. 133	227a			3
Ex. 134	227b			3
Ex. 135	229a			1
Ex. 136	229b			0.04
Ex. 137	230a			5
Ex. 138	230b			12

Ex. 139	234			1
Ex. 140	235			0.004
Ex. 141	238a		0% @ 1 μ M	0.05
Ex. 142	238b		20% @ 1 μ M	0.76
Ex. 143	239a		0%	11.6
Ex. 144	239b		14%	8.1
Ex. 145	243a		6%	1.6
Ex. 146	243b			99
Ex. 147	243c			4
Ex. 148	243d			3

Ex. 149	243e			12
Ex. 150	243f			3
Ex. 151	243g			3
Ex. 152	243h			3
Ex. 153	246a			5
Ex. 154	246b			14
Ex. 155	248			4
Ex. 156	251a		0%	0.7
Ex. 157	251b			1
Ex. 158	251c			2

	252		0%	> 20
	253		0%	> 20

* Unless otherwise indicated, inhibition values expressed in percent refer to the percent inhibition at 20 μ M.

Example 160:

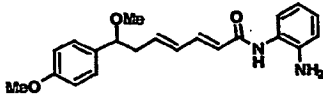
5 Inhibition of Histone Deacetylase in Whole Cells

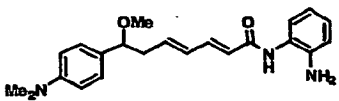
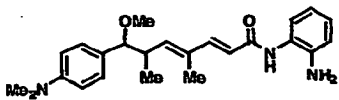
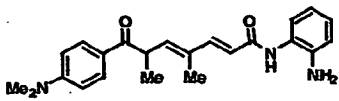
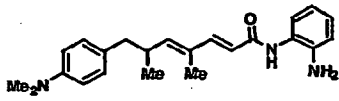
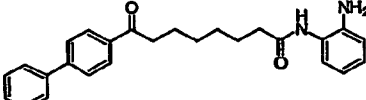
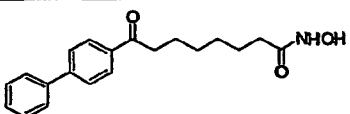
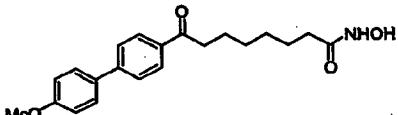
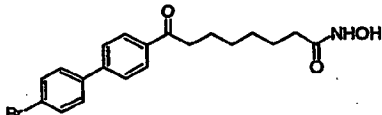
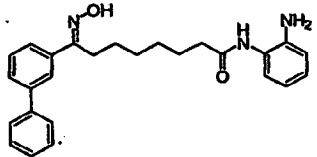
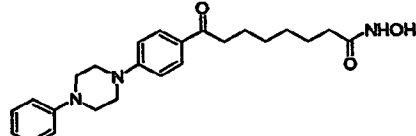
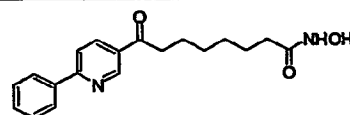
1. Histone H4 acetylation in whole cells by immunoblots

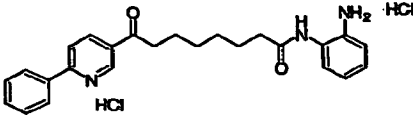
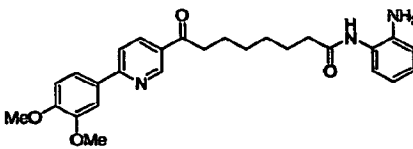
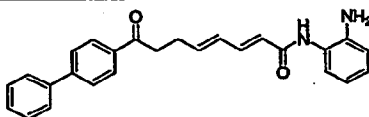
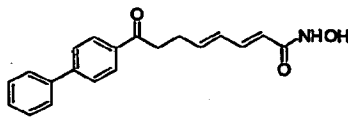
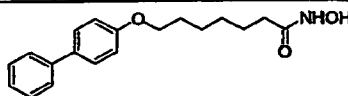
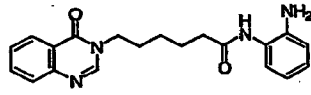
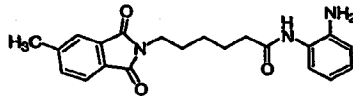
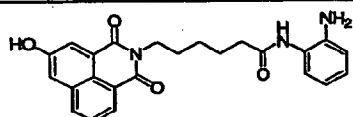
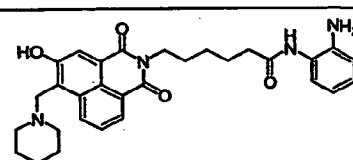
T24 human bladder cancer cells growing in culture were incubated with HDAC inhibitors for 16 hours. Histones were extracted from the cells after the culture period as described by M. Yoshida *et al.* (*J. Biol. Chem.* 265(28): 17174-17179 (1990)). 20 μ g of total histone protein was loaded onto SDS/PAGE and transferred to nitrocellulose membranes. Membranes were probed with polyclonal antibodies specific for acetylated histone H-4 (Upstate Biotech Inc.), followed by horse radish peroxidase conjugated secondary antibodies (Sigma). Enhanced Chemiluminescence (ECL) (Amersham) detection was performed using Kodak films (Eastman Kodak). Acetylated H-4 signal was quantified by densitometry.

Data for selected compounds are presented in Table 3. Data are presented as the concentration effective for reducing the acetylated H-4 signal by 50% (EC_{50}).

Table 3: Inhibition of Histone Acetylation in Cells

Example	Cpd.	Structure	EC_{50} (μ M)
Ex. 4	18		10

Ex. 6	20		5
Ex. 12	34		5
Ex. 13	35		5
Ex. 15	39		10
Ex. 32	77		10
Ex. 33	78		10
Ex. 69	120		10
Ex. 70	124		3
Ex. 78	134		10
Ex. 86	138g		5
Ex. 91	145		10

Ex. 92	146		5
Ex. 94	151		3
Ex. 120	204		15
Ex. 121	205		3
Ex. 140	235		5
Ex. 143	239a		5
Ex. 148	243d		3
Ex. 156	251a		5
Ex. 158	251c		1

2. Acid Urea Triton (AUT) gel analysis of histone acetylation.

Human cancer cells (T24, 293T or Jurkat cells) growing in culture are incubated with HDAC inhibitors for 24 h. Histones are extracted from the cells as described by M. Yoshida *et al.* (*J. Biol. Chem.* 265(28): 17174-17179 (1990)). Acid urea triton (AUT) gel electrophoresis is used for detection of acetylated histone m

olecules. Histones (150 µg of total protein) are electrophoresed at 80 V for 16 h at room temperature as described by M. Yoshida *et al.*, *supra*. Gels are stained with Coomassie brilliant blue to visualize histones, dried and scanned by densitometry to quantified acetylation of histones.

5

Example 161:**Antineoplastic Effect of Histone Deacetylase Inhibitors on Tumor Cells *In Vivo***

Eight to ten week old female BALB/c nude mice (Taconic Labs, Great Barrington, NY) were injected subcutaneously in the flank area with 2×10^6 preconditioned A549 human lung carcinoma cells. Preconditioning of these cells was accomplished by a minimum of three consecutive tumor transplantations in the same strain of nude mice. Subsequently, tumor fragments of approximately 30 mgs were excised and implanted subcutaneously in mice, in the left flank area, under isoflurane anesthesia (Abbott Labs, Geneva, Switzerland). When the tumors reached a mean volume of 100 mm^3 , the mice were treated intravenously, subcutaneously or intraperitoneally, by daily injection of a solution of the inhibitor in an appropriate vehicle such as PBS, DMSO/water, or Tween 80/water, at a dose from about 10 mg/kg to about 50 mg/kg, for 21 days. Tumor volume was calculated every second day post infusion according to standard methods (*e.g.*, Meyer *et al.*, *Int. J. Cancer* 43: 851-856 (1989)). Treatment with compound 34 caused a significant reduction in tumor weight and volume relative to controls treated with saline only (*i.e.*, no HDAC inhibitor). In addition, the activity of histone deacetylase, when measured, is expected to be significantly reduced relative to saline treated controls.

Example 162:**Synergistic Antineoplastic Effect of Histone Deacetylase Inhibitors and Histone Deacetylase Antisense Oligonucleotides on Tumor Cells *In Vivo***

The purpose of this example is to illustrate the ability of the histone deacetylase inhibitor of the invention and a histone deacetylase antisense

oligonucleotide to synergistically inhibit tumor growth in a mammal. Preferably, the antisense oligonucleotide and the HDAC inhibitor inhibit the expression and activity of the same histone deacetylase.

5 As described in Example 161, mice bearing implanted A549 tumors (mean volume 100 mm^3) are treated daily with saline preparations containing from about 0.1 mg to about 30 mg per kg body weight of histone deacetylase antisense oligonucleotide. A second group of mice is treated daily with pharmaceutically acceptable preparations containing from about 0.01 mg to about 5 mg per kg body weight of HDAC inhibitor.

10 Some mice receive both the antisense oligonucleotide and the HDAC inhibitor. Of these mice, one group may receive the antisense oligonucleotide and the HDAC inhibitor simultaneously intravenously via the tail vein. Another group may receive the antisense oligonucleotide via the tail vein, and the HDAC inhibitor subcutaneously. Yet another group may receive both the antisense
15 oligonucleotide and the HDAC inhibitor subcutaneously. Control groups of mice are similarly established which receive no treatment (*e.g.*, saline only), a mismatch antisense oligonucleotide only, a control compound that does not inhibit histone deacetylase activity, and a mismatch antisense oligonucleotide with a control compound.

20 Tumor volume is measured with calipers. Treatment with the antisense oligonucleotide plus the histone deacetylase protein inhibitor according to the invention causes a significant reduction in tumor weight and volume relative to controls.

What is claimed is:

1. An inhibitor of histone deacetylase represented by formula (1):



wherein

- 5 Cy is cycloalkyl, aryl, or a radical of a heterocyclic moiety, any of which may be optionally substituted;

X is selected from the group consisting of C=O, C=CH₂, CH(OH), CH(OR¹), C=N(OH), and C=N(OR¹), where R¹ is alkyl, aryl, aralkyl, or acyl;

- 10 Y¹ is a C₃-C₇ alkylene, wherein said alkylene may be optionally substituted, and wherein one or two carbon atoms in the linear chain connecting X and W may be replaced with O, NR³, or S(O)_n, where R³ is hydrogen, alkyl, aryl, aralkyl, sulfonyl, acyl, alkoxycarbonyl, or carbamoyl, and n is 0, 1, or 2, provided that the atoms in Y¹ that are attached to X and to W are carbon atoms, and further provided that Y¹ does not comprise an ester or amide linkage in the linear chain
15 connecting X and W; and

W is selected from the group consisting of -C(O)-CH₂-SR², -C(O)-NH-OM, -NH-C(O)-NH-Z, and -C(O)-NH-Z, where

R² is alkyl, aryl, aralkyl, or acyl, wherein the aryl portion of any such groups may be optionally substituted;

- 20 M is hydrogen or a pharmaceutically acceptable cation;

Z is selected from the group consisting of anilinyll, pyridyl, thiazolyl, hydroxyphenyl, thiadiazolyl, anilinylmethyl, or pyridylmethyl, any of which groups optionally may be substituted with halo, hydroxy, amino, nitro, C₁-C₄ alkyl, or C₁-C₄ alkoxy;

- 25 provided that X is C=CH₂, CH(OR¹), C=N(OH), or C=N(OR¹) when W is -C(O)-NH-OM and Cy is unsubstituted phenyl, dimethylaminophenyl, or methoxyphenyl; and

further provided that when W is $-C(O)-CH_2-SR^2$, the carbon atom in Y^1 that is attached to W is unsubstituted or is substituted with other than amino, acylamino, alkoxycarbonyl, or carbamoyl.

2. The inhibitor of claim 1, wherein Cy is C_6-C_{10} aryl or is a radical of a heterocyclic moiety selected from the group consisting of thiophene, benzothiophene, furan, benzofuran, pyridine, quinoline, indole, isoquinoline, thiazole, morpholine, piperidine, and piperazine, any of which groups may be optionally substituted.

3. The inhibitor of claim 2, wherein the aryl or heterocyclic moiety is substituted by one or two substituents independently selected from the group consisting of C_1-C_4 alkyl, C_1-C_4 haloalkyl, C_6-C_{10} aryl, heteroaryl, heterocyclyl, $(C_6-C_{10})ar(C_1-C_6)alkyl$, halo, nitro, hydroxy, C_1-C_6 alkoxy, C_6-C_{10} aryloxy, heteroaryloxy, C_1-C_6 alkoxycarbonyl, C_6-C_{10} aryloxycarbonyl, heteroaryloxycarbonyl, carboxy, and amino.

4. The inhibitor of claim 1, wherein Cy has the formula $-Cy^1-Cy^2$ or $-Cy^1-G-Cy^2$, wherein Cy^1 and Cy^2 are independently C_3-C_6 cycloalkyl, C_6-C_{10} aryl, or a radical of a heterocyclic moiety, which groups optionally may be substituted, and G is O, NR^3 , or $S(O)_n$, where R^3 is hydrogen, alkyl, aryl, aralkyl, sulfonyl, acyl, alkoxycarbonyl, or carbamoyl, and n is 0, 1, or 2.

5. The inhibitor of claim 4, wherein Cy^1 and Cy^2 are independently selected from the group consisting of phenyl, pyridinyl, morpholinyl, piperidinyl, piperazinyl, which groups optionally may be substituted.

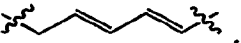
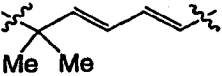
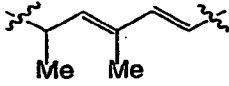
6. The inhibitor of claim 1, wherein X is selected from the group consisting of $CH(OR^1)$, $C=N(OH)$, and $C=N(OR^1)$, where R^1 is C_1-C_6 alkyl, C_6-C_{10} aryl, or $(C_6-C_{10})ar(C_1-C_6)alkyl$.

7. The inhibitor of claim 1, wherein one to about three carbon atoms of the alkylene are independently substituted with halo, oxo, oximino, nitro, haloalkyl, alkyl, aralkyl, alkoxy, aryloxy, alkoxycarbonyl, carboxy, hydroxyalkyl, acyl, acyloxy, or cyano.

5 8. The inhibitor of claim 1, wherein Y^1 comprises an all-carbon linear chain connecting X and W.

9. The inhibitor of claim 8, wherein the linear chain connecting X and W comprises a dienyl moiety, wherein the dienyl moiety is attached to W.

10. The inhibitor of claim 9, wherein Y^1 is selected from the group

10 consisting of , , and 

11. The inhibitor of claim 8, wherein Y^1 is $-(CH_2)_m$, where m is 5, 6, or 7.

12. The inhibitor of claim 1, wherein one carbon atom in the linear chain connecting X and W is replaced with O, NR^3 , or $S(O)_n$.

13. The inhibitor of claim 12, wherein Y^1 is $-(CH_2)_n-S(O)_n-(CH_2)_p$, where n is 0, 1, or 2, and p is 3, 4, or 5.

14. The inhibitor of claim 1, wherein W is $-C(O)-NH-OM$, M being selected from the group consisting of hydrogen, sodium, potassium, magnesium, and calcium.

15 15. The inhibitor of claim 1, wherein W is $-C(O)-NH-Z$ or $-NH-C(O)-NH-Z$, Z being unsubstituted 2-anilinyI or unsubstituted 2-pyridyl.

16. The inhibitor of claim 1, wherein W is $-C(O)-CH_2-SR^2$, R^2 being selected from the group consisting of C_1-C_6 alkyl, C_6-C_{10} aryl, $(C_6-C_{10})ar(C_1-C_6)alkyl$,

(C₁-C₆ alkyl)carbonyl, (C₆-C₁₀ aryl)carbonyl, and ((C₆-C₁₀)ar(C₁-C₆)alkyl)carbonyl, wherein the aryl portion of any such groups may be optionally substituted.

17. The inhibitor of claim 16, wherein R² is selected from the group consisting of methyl, phenyl, benzyl, benzoyl, and acetyl.

5 18. An inhibitor of histone deacetylase represented by formula (2):



wherein

Cy is cycloalkyl, aryl, or a radical of a heterocyclic moiety, any of which may be optionally substituted;

10 Y² is C₅-C₇ alkylene, wherein said alkylene may be optionally substituted, and wherein one or two carbon atoms in the linear chain connecting Cy and W may be replaced with O, NR³, or S(O)_n, where R³ is hydrogen, alkyl, aryl, aralkyl, sulfonyl, acyl, alkoxycarbonyl, or carbamoyl, and n is 0, 1, or 2, provided that Y² does not comprise an ester or amide linkage in the linear chain connecting Cy and

15 W; and

W is selected from the group consisting of -C(O)-CH₂-SR², -NH-C(O)-NH-Z, and -C(O)-NH-Z, where

R² is alkyl, aryl, aralkyl, or acyl, wherein the aryl portion of any such groups may be optionally substituted; and

20 Z is selected from the group consisting of anilinyll, pyridyl, thiazolyl, hydroxyphenyl, thiadiazolyl, anilinyllmethyl, or pyridylmethyl, any of which groups optionally may be substituted with halo, hydroxy, amino, nitro, C₁-C₄ alkyl, or C₁-C₄ alkoxy;

provided that when W is -C(O)-CH₂-SR², the carbon atom in Y² that is attached to
25 W is unsubstituted or is substituted with other than amino, acylamino, alkoxycarbonyl, or carbamoyl.

19. The inhibitor of claim 18, wherein Cy is C₆-C₁₀ aryl or is a radical of a heterocyclic moiety selected from the group consisting of thiophene, benzothiophene, furan, benzofuran, pyridine, quinoline, indole, isoquinoline, thiazole, morpholine, piperidine, piperazine, quinazolinone, benzotriazinone, phthalimide, and dioxobenzoisoquinoline, any of which groups may be optionally substituted.

20. The inhibitor of claim 18, wherein the aryl or heterocyclic moiety is substituted by one or two substituents independently selected from the group consisting of C₁-C₄ alkyl, C₁-C₄ haloalkyl, C₆-C₁₀ aryl, heteroaryl, heterocyclyl, (C₆-C₁₀)ar(C₁-C₆)alkyl, halo, nitro, hydroxy, C₁-C₆ alkoxy, C₆-C₁₀ aryloxy, heteroaryloxy, C₁-C₆ alkoxycarbonyl, C₆-C₁₀ aryloxycarbonyl, heteroaryloxycarbonyl, carboxy, and amino.

21. The inhibitor of claim 20, wherein Cy has the formula -Cy¹-Cy² or -Cy¹-G-Cy², wherein Cy¹ and Cy² are independently C₃-C₆ cycloalkyl, C₆-C₁₀ aryl, or a radical of a heterocyclic moiety, which groups optionally may be substituted, and G is O, NR³, or S(O)_n, where R³ is hydrogen, alkyl, aryl, aralkyl, sulfonyl, acyl, alkoxycarbonyl, or carbamoyl, and n is 0, 1, or 2.

22. The inhibitor of claim 21, wherein Cy¹ and Cy² are independently selected from the group consisting of phenyl, pyridinyl, morpholinyl, piperidinyl, piperazinyl, which groups optionally may be substituted.

23. The inhibitor of claim 18, wherein one to about four carbon atoms of the alkylene are independently substituted with halo, oxo, oximino, nitro, haloalkyl, alkyl, aralkyl, alkoxy, aryloxy, alkoxycarbonyl, carboxy, hydroxyalkyl, acyl, acyloxy, or cyano.

24. The inhibitor of claim 18, wherein one carbon atom in the linear chain connecting Cy and W is replaced with O, NR³, or S(O)_n.

25. The inhibitor of claim 19, wherein one carbon atom in the linear chain connecting Cy and W is replaced with NR^3 , where R^3 is selected from the group consisting of $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_6\text{-C}_{10}$ aryl, $(\text{C}_6\text{-C}_{10})\text{ar}(\text{C}_1\text{-C}_6)\text{alkyl}$, $(\text{C}_1\text{-C}_6\text{alkyl})\text{oxycarbonyl}$, $(\text{C}_6\text{-C}_{10}\text{aryl})\text{oxycarbonyl}$, $((\text{C}_6\text{-C}_{10})\text{ar}(\text{C}_1\text{-C}_6)\text{alkyl})\text{oxycarbonyl}$,
 5 $(\text{C}_1\text{-C}_6\text{alkyl})\text{carbonyl}$, $(\text{C}_6\text{-C}_{10}\text{aryl})\text{carbonyl}$, and $((\text{C}_6\text{-C}_{10})\text{ar}(\text{C}_1\text{-C}_6)\text{alkyl})\text{carbonyl}$.

26. The inhibitor of claim 18, wherein one or two carbon atoms in the linear chain connecting Cy and W are replaced by O.

27. The inhibitor of claim 18, wherein W is $-\text{C}(\text{O})\text{-NH-Z}$ or $-\text{NH-C}(\text{O})\text{-NH-Z}$, Z being unsubstituted 2-anilinyI or unsubstituted 2-pyridyl.

10 28. The inhibitor of claim 18, wherein W is $-\text{C}(\text{O})\text{-CH}_2\text{-SR}^2$, R^2 being selected from the group consisting of $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_6\text{-C}_{10}$ aryl, $(\text{C}_6\text{-C}_{10})\text{ar}(\text{C}_1\text{-C}_6)\text{alkyl}$, $(\text{C}_1\text{-C}_6\text{alkyl})\text{carbonyl}$, $(\text{C}_6\text{-C}_{10}\text{aryl})\text{carbonyl}$, and $((\text{C}_6\text{-C}_{10})\text{ar}(\text{C}_1\text{-C}_6)\text{alkyl})\text{carbonyl}$, wherein the aryl portion of any such groups may be optionally substituted.

15 29. The inhibitor of claim 28, wherein R^2 is selected from the group consisting of methyl, phenyl, benzyl, benzoyl, and acetyl.

30. An inhibitor of histone deacetylase represented by formula (3):



wherein

20 Cy is cycloalkyl, aryl, or a radical of a heterocyclic moiety, any of which may be optionally substituted, provided that Cy is other than dimethylaminonaphthyl when Y^3 is $-(\text{CH}_2)_3\text{-}$;

Y^3 is $\text{C}_2\text{-C}_6$ alkylene, wherein said alkylene may be optionally substituted with one or more substituents independently selected from the group consisting of halo, hydroxy, oxo, nitro, haloalkyl, alkyl, aralkyl, alkoxy, aryloxy, carboxy,
 25 hydroxyalkyl, acyl, acyloxy, and cyano; and

W is selected from the group consisting of $-C(O)-CH_2-SR^2$, $-C(O)-NH-OM$, $-NH-C(O)-NH-Z$, and $-C(O)-NH-Z$, where

R^2 is alkyl, aryl, aralkyl, or acyl, wherein the aryl portion of any such groups may be optionally substituted;

5 M is hydrogen or a pharmaceutically acceptable cation; and

Z is selected from the group consisting of aniliny, pyridyl, thiazolyl, hydroxyphenyl, thiadiazolyl, anilinylmethyl, or pyridylmethyl, any of which groups optionally may be substituted with halo, hydroxy, amino, nitro, C_1-C_4 alkyl, or C_1-C_4 alkoxy;

10 provided that Z does not have the formula $-(C_5H_5N)-NHC(O)-Y^3-NH-S(O)_2-Cy$.

31. The inhibitor of claim 30, wherein Cy is C_6-C_{10} aryl or is a radical of a heterocyclic moiety selected from the group consisting of thiophene, benzothiophene, furan, benzofuran, pyridine, quinoline, indole, isoquinoline, thiazole, morpholine, piperidine, and piperazine, any of which groups may be
15 optionally substituted.

32. The inhibitor of claim 31, wherein the aryl or heterocyclic moiety is substituted by one or two substituents independently selected from the group consisting of C_1-C_4 alkyl, C_1-C_4 haloalkyl, C_6-C_{10} aryl, heteroaryl, heterocyclyl, $(C_6-C_{10})ar(C_1-C_6)alkyl$, halo, nitro, hydroxy, C_1-C_6 alkoxy, C_6-C_{10} aryloxy, heteroaryloxy,
20 C_1-C_6 alkoxycarbonyl, C_6-C_{10} aryloxycarbonyl, heteroaryloxycarbonyl, carboxy, and amino.

33. The inhibitor of claim 30, wherein Cy has the formula $-Cy^1-Cy^2$ or $-Cy^1-G-Cy^2$, wherein Cy^1 and Cy^2 are independently C_3-C_6 cycloalkyl, C_6-C_{10} aryl, or a radical of a heterocyclic moiety, which groups optionally may be substituted,
25 and G is O, NR^3 , or $S(O)_n$, where R^3 is hydrogen, alkyl, aryl, aralkyl, sulfonyl, acyl, alkoxycarbonyl, or carbamoyl, and n is 0, 1, or 2.

34. The inhibitor of claim 33, wherein Cy^1 and Cy^2 are independently selected from the group consisting of phenyl, pyridinyl, morpholinyl, piperidinyl, piperazinyl, which groups optionally may be substituted.

5 35. The inhibitor of claim 30, wherein Y^3 is a C_2-C_6 alkylene optionally substituted with one or two non-hydrogen substituents independently selected from the group consisting of halo, hydroxy, oxo, nitro, (halo)₁₋₅(C_1-C_3)alkyl, C_1-C_6 alkyl, (C_6-C_{10})ar(C_1-C_6)alkyl, C_1-C_6 alkoxy, C_6-C_{10} aryloxy, carboxy, hydroxy(C_1-C_6)alkyl, C_1-C_6 alkylcarbonyl, C_6-C_{10} arylcarbonyl, C_1-C_6 alkylcarbonyloxy, C_6-C_{10} arylcarbonyloxy, and cyano.

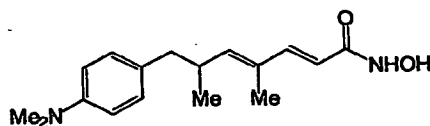
10 36. The inhibitor of claim 33, wherein Y^3 is an optionally substituted saturated C_4-C_5 alkylene.

37. The inhibitor of claim 30, wherein W is $-C(O)-NH-OM$, M being selected from the group consisting of hydrogen, sodium, potassium, magnesium, and calcium.

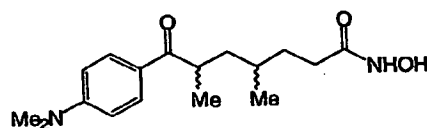
15 38. The inhibitor of claim 30, wherein W is $-C(O)-NH-Z$ or $-NH-C(O)-NH-Z$, Z being unsubstituted 2-anilinyl or unsubstituted 2-pyridyl.

39. The inhibitor of claim 30, wherein W is $-C(O)-CH_2-SR^2$, R^2 being selected from the group consisting of C_1-C_6 alkyl, C_6-C_{10} aryl, (C_6-C_{10})ar(C_1-C_6)alkyl, (C_1-C_6 alkyl)carbonyl, (C_6-C_{10} aryl)carbonyl, and ((C_6-C_{10})ar(C_1-C_6)alkyl)carbonyl, wherein the aryl portion of any such groups may be optionally substituted.

20 40. An inhibitor of histone deacetylase represented by one of formulae (4)-(5):

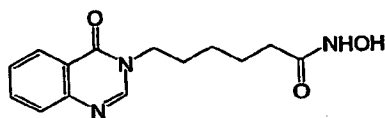


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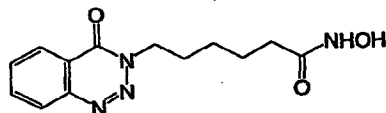
(5)

41. An inhibitor of histone deacetylase represented by one of formulae (6)-(7):



5

(6)



(7)

42. A pharmaceutical composition comprising an inhibitor of histone deacetylase represented by formula (1):

10



wherein

Cy is cycloalkyl, aryl, or a radical of a heterocyclic moiety, any of which may be optionally substituted;

- 15 X is selected from the group consisting of C=O, C=CH₂, CH(OH), CH(OR¹), C=N(OH), and C=N(OR¹), where R¹ is alkyl, aryl, aralkyl, or acyl;

- 20 Y¹ is a C₃-C₇ alkylene, wherein said alkylene may be optionally substituted, and wherein one or two carbon atoms in the linear chain connecting X and W may be replaced with O, NR³, or S(O)_n, where R³ is hydrogen, alkyl, aryl, aralkyl, sulfonyl, acyl, alkoxy carbonyl, or carbamoyl, and n is 0, 1, or 2, provided that the atoms in Y¹ that are attached to X and to W are carbon atoms, and further provided that Y¹ does not comprise an ester or amide linkage in the linear chain connecting X and W; and

W is selected from the group consisting of $-C(O)-CH_2-SR^2$, $-C(O)-NH-OM$, $-NH-C(O)-NH-Z$, and $-C(O)-NH-Z$, where

R^2 is alkyl, aryl, aralkyl, or acyl, wherein the aryl portion of any such groups may be optionally substituted;

5 M is hydrogen or a pharmaceutically acceptable cation;

Z is selected from the group consisting of aniliny, pyridyl, thiazolyl, hydroxyphenyl, thiadiazolyl, anilinylmethyl, or pyridylmethyl, any of which groups optionally may be substituted with halo, hydroxy, amino, nitro, C_1-C_4 alkyl, or C_1-C_4 alkoxy; and

10 a pharmaceutically acceptable carrier, excipient, or diluent;

provided that X is $C=CH_2$, $CH(OR^1)$, $C=N(OH)$, or $C=N(OR^1)$ when W is $-C(O)-NH-OM$ and Cy is unsubstituted phenyl, dimethylaminophenyl, or methoxyphenyl; and

further provided that when W is $-C(O)-CH_2-SR^2$, the carbon atom in Y^1 that is
15 attached to W is unsubstituted or is substituted with other than amino, acylamino, alkoxycarbonyl, or carbamoyl.

43. A pharmaceutical composition comprising an inhibitor of histone deacetylase represented by formula (2):



20 wherein

Cy is cycloalkyl, aryl, or a radical of a heterocyclic moiety, any of which may be optionally substituted;

Y^2 is C_5-C_7 alkylene, wherein said alkylene may be optionally substituted, and wherein one or two carbon atoms in the linear chain connecting Cy and W
25 may be replaced with O, NR^3 , or $S(O)_n$, where R^3 is hydrogen, alkyl, aryl, aralkyl, sulfonyl, acyl, alkoxycarbonyl, or carbamoyl, and n is 0, 1, or 2, provided that Y^2 does not comprise an ester or amide linkage in the linear chain connecting Cy and W; and

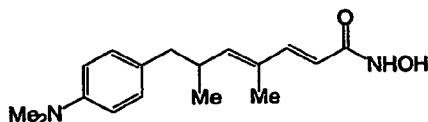
M is hydrogen or a pharmaceutically acceptable cation; and

Z is selected from the group consisting of anilinyll, pyridyl, thiazolyl, hydroxyphenyl, thiadiazolyl, anilinylmethyl, or pyridylmethyl, any of which groups optionally may be substituted with halo, hydroxy, amino, nitro, C₁-C₄ alkyl, or C₁-C₄ alkoxy; and

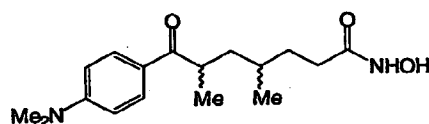
a pharmaceutically acceptable carrier, excipient, or diluent.

provided that Z does not have the formula $-(C_3H_5N)-NHC(O)-Y^2-NH-S(O)_2-Cy$.

45. A pharmaceutical composition comprising an inhibitor of histone deacetylase represented by one of formulae (4)-(5):



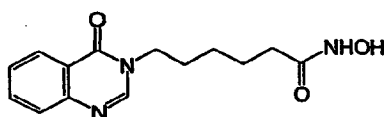
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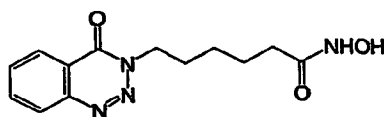
(5);

and a pharmaceutically acceptable carrier, excipient, or diluent.

46. A pharmaceutical composition comprising an inhibitor of histone deacetylase represented by one of formulae (6)-(7):



(6)



(7);

and a pharmaceutically acceptable carrier, excipient, or diluent.

47. A method of inhibiting histone deacetylase in a cell, comprising contacting a cell in which inhibition of histone deacetylase is desired with an inhibitor of histone deacetylase represented by formula (1):



5 wherein

Cy is cycloalkyl, aryl, or a radical of a heterocyclic moiety, any of which may be optionally substituted;

X is selected from the group consisting of C=O, C=CH₂, CH(OH), CH(OR¹), C=N(OH), and C=N(OR¹), where R¹ is alkyl, aryl, aralkyl, or acyl;

10 Y¹ is a C₃-C₇ alkylene, wherein said alkylene may be optionally substituted, and wherein one or two carbon atoms in the linear chain connecting X and W may be replaced with O, NR³, or S(O)_n, where R³ is hydrogen, alkyl, aryl, aralkyl, sulfonyl, acyl, alkoxy carbonyl, or carbamoyl, and n is 0, 1, or 2, provided that the atoms in Y¹ that are attached to X and to W are carbon atoms, and further
15 provided that Y¹ does not comprise an ester or amide linkage in the linear chain connecting X and W; and

W is selected from the group consisting of -C(O)-CH₂-SR², -C(O)-NH-OM, -NH-C(O)-NH-Z, and -C(O)-NH-Z, where

20 R² is alkyl, aryl, aralkyl, or acyl, wherein the aryl portion of any such groups may be optionally substituted;

M is hydrogen or a pharmaceutically acceptable cation;

Z is selected from the group consisting of anilinyll, pyridyl, thiazolyl, hydroxyphenyl, thiadiazolyl, anilinyllmethyl, or pyridylmethyl, any of which groups optionally may be substituted with halo, hydroxy, amino,
25 nitro, C₁-C₄ alkyl, or C₁-C₄ alkoxy;

provided that X is C=CH₂, CH(OR¹), C=N(OH), or C=N(OR¹) when W is -C(O)-NH-OM and Cy is unsubstituted phenyl, dimethylaminophenyl, or methoxyphenyl; and

further provided that when W is $-C(O)-CH_2-SR^2$, the carbon atom in Y^1 that is attached to W is unsubstituted or is substituted with other than amino, acylamino, alkoxycarbonyl, or carbamoyl.

48. A method of inhibiting histone deacetylase in a cell, comprising
5 contacting a cell in which inhibition of histone deacetylase is desired with an inhibitor of histone deacetylase represented by formula (2):



wherein

- 10 Cy is cycloalkyl, aryl, or a radical of a heterocyclic moiety, any of which may be optionally substituted;

- Y^2 is C_5-C_n alkylene, wherein said alkylene may be optionally substituted, and wherein one or two carbon atoms in the linear chain connecting Cy and W may be replaced with O, NR^3 , or $S(O)_n$, where R^3 is hydrogen, alkyl, aryl, aralkyl, sulfonyl, acyl, alkoxycarbonyl, or carbamoyl, and n is 0, 1, or 2, provided that Y^2
15 does not comprise an ester or amide linkage in the linear chain connecting Cy and W; and

W is selected from the group consisting of $-C(O)-CH_2-SR^2$, $-NH-C(O)-NH-Z$, and $-C(O)-NH-Z$, where

- 20 R^2 is alkyl, aryl, aralkyl, or acyl, wherein the aryl portion of any such groups may be optionally substituted; and

Z is selected from the group consisting of anilinyll, pyridyl, thiazolyl, hydroxyphenyl, thiadiazolyl, anilinylmethyl, or pyridylmethyl, any of which groups optionally may be substituted with halo, hydroxy, amino, nitro, C_1-C_4 alkyl, or C_1-C_4 alkoxy;

- 25 provided that when W is $-C(O)-CH_2-SR^2$, the carbon atom in Y^2 that is attached to W is unsubstituted or is substituted with other than amino, acylamino, alkoxycarbonyl, or carbamoyl.

49. A method of inhibiting histone deacetylase in a cell, comprising contacting a cell in which inhibition of histone deacetylase is desired with an inhibitor of histone deacetylase represented by formula (3):



5 wherein

Cy is cycloalkyl, aryl, or a radical of a heterocyclic moiety, any of which may be optionally substituted, provided that Cy is other than dimethylaminonaphthyl when Y³ is -(CH₂)₃-;

10 Y³ is C₂-C₆ alkylene, wherein said alkylene may be optionally substituted with one or more substituents independently selected from the group consisting of halo, hydroxy, oxo, nitro, haloalkyl, alkyl, aralkyl, alkoxy, aryloxy, carboxy, hydroxyalkyl, acyl, acyloxy, and cyano; and

W is selected from the group consisting of -C(O)-CH₂-SR², -C(O)-NH-OM, -NH-C(O)-NH-Z, and -C(O)-NH-Z, where

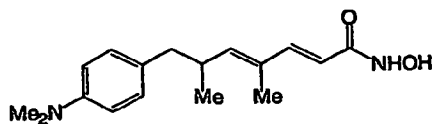
15 R² is alkyl, aryl, aralkyl, or acyl, wherein the aryl portion of any such groups may be optionally substituted;

M is hydrogen or a pharmaceutically acceptable cation; and

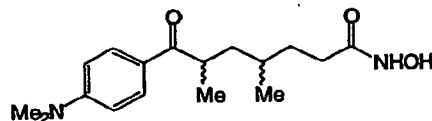
20 Z is selected from the group consisting of aniliny, pyridyl, thiazolyl, hydroxyphenyl, thiadiazolyl, anilinylmethyl, or pyridylmethyl, any of which groups optionally may be substituted with halo, hydroxy, amino, nitro, C₁-C₄ alkyl, or C₁-C₄ alkoxy;

provided that Z does not have the formula -(C₅H₃N)-NHC(O)-Y³-NH-S(O)₂-Cy.

50. A method of inhibiting histone deacetylase in a cell, comprising contacting a cell in which inhibition of histone deacetylase is desired with an
25 inhibitor of histone deacetylase represented by one of formulae (4)-(5):

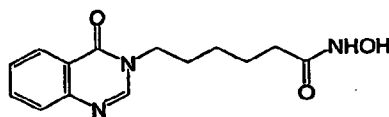


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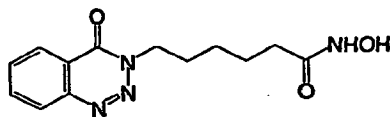


(5)

- 5 51. A method of inhibiting histone deacetylase in a cell, comprising contacting a cell in which inhibition of histone deacetylase is desired with an inhibitor of histone deacetylase represented by one of formulae (6)-(7):



(6)



(7)

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52. The method of any one of claims 47-51, wherein cell proliferation is inhibited in the contacted cell.

- 15 53. The method of any one of claims 47-51, wherein the cell is a neoplastic cell.

54. The method of claim 53, wherein the neoplastic cell is in an animal.

55. The method of claim 54, wherein the neoplastic cell is in a neoplastic growth.

56. The method of any one of claims 47-51, further comprising contacting the cell with an antisense oligonucleotide that inhibits the expression of a histone deacetylase.